

Meet The Professor

Optimizing the Selection and Sequencing of Therapy for Patients with HER2-Positive Breast Cancer

Adam M Brufsky, MD, PhD

Professor of Medicine

Co-Director, Comprehensive Breast Cancer Center

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Associate Division Chief, Division of Hematology/Oncology

Department of Medicine

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Commercial Support

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Dr Love — Disclosures

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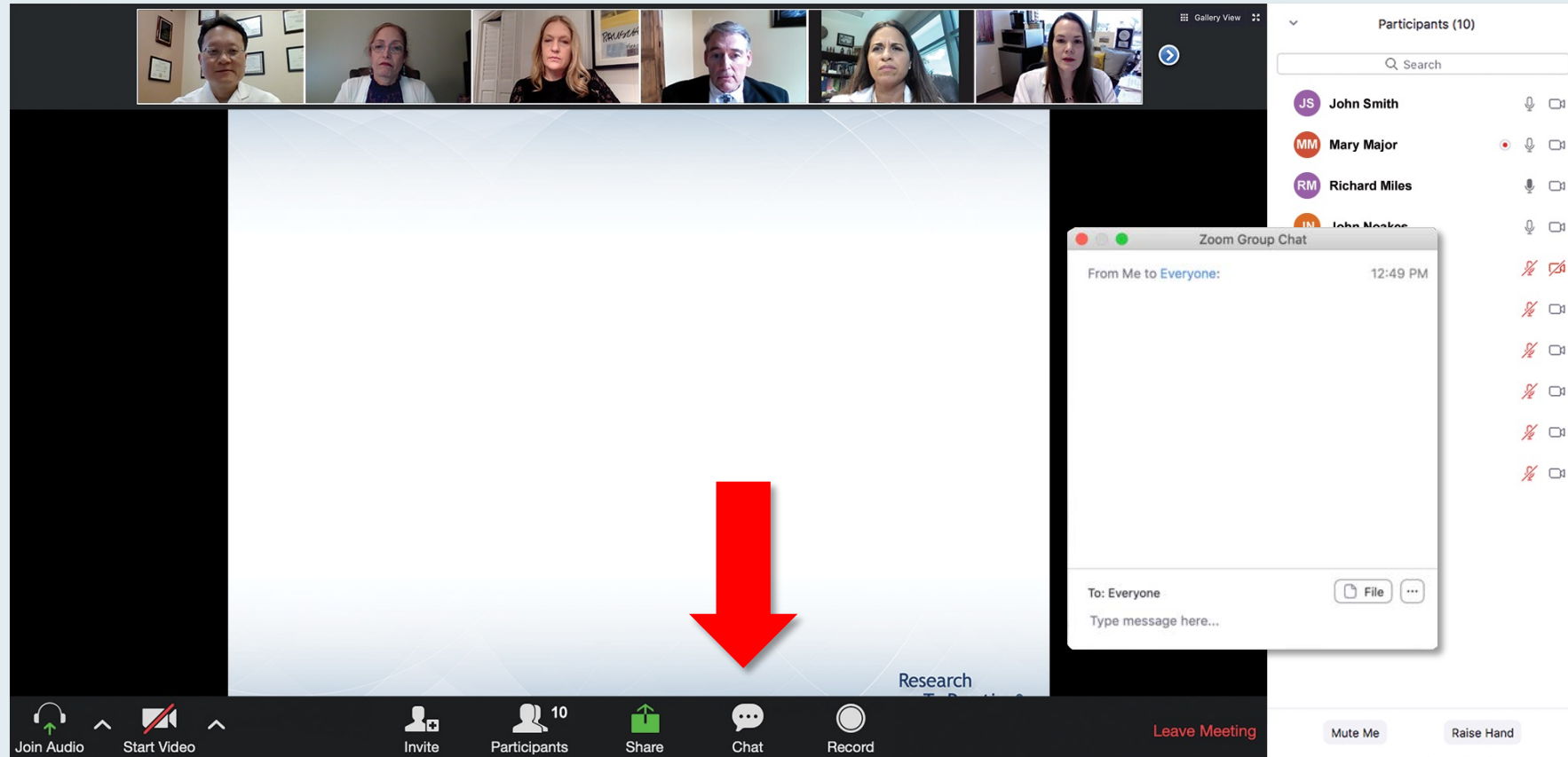
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Dr Brufsky — Disclosures

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We Encourage Clinicians in Practice to Submit Questions



Feel free to submit questions now before the program begins and throughout the program.

Familiarizing Yourself with the Zoom Interface

Expand chat submission box

The screenshot shows a Zoom meeting interface. At the top, there are video thumbnails for participants: RTP Coordinat..., Kirsten Miller, RTP Mike Rivera, and Lisa Suarez. Below the thumbnails is a slide titled "Meet The Professor Program Steering Committee" with six members listed:

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Assistant Professor of Medicine
Weill Cornell Medicine
New York, New York
- Ian W Flinn, MD, PhD**
Director of Lymphoma Research Program
Sarah Cannon Research Institute
Tennessee Oncology
Nashville, Tennessee
- Steven Coutre, MD**
Professor of Medicine (Hematology)
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Stanford, California
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Harvard Medical School
Director of Clinical Research
Division of Lymphoma
Dana-Farber Cancer Institute
Boston, Massachusetts
- Brian T Hill, MD, PhD**
Director, Lymphoid Malignancy Program
Cleveland Clinic Taussig Cancer Institute
Cleveland, Ohio

On the right side, there is a chat window titled "Chat". It shows two messages from "Me to Panelists" and "Me to Panelists and Attendees" at 4:31 PM and 4:32 PM respectively. Each message contains a welcome message and a link to a PDF file: http://images.researchtopractice.com/2021/Meetings/Slides/MTP_ToGo_CLL_2021_April1.pdf. Below the messages is a "To:" dropdown menu set to "Panelists and Attendees" and a text input field labeled "Type message here...". A red arrow points to the white line above the text input field, indicating where to drag to expand the box.

Drag the white line above the submission box up to create more space for your message.

Familiarizing Yourself with the Zoom Interface

Increase chat font size



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**Press Command (for Mac) or Control (for PC) and the + symbol.
You may do this as many times as you need for readability.**

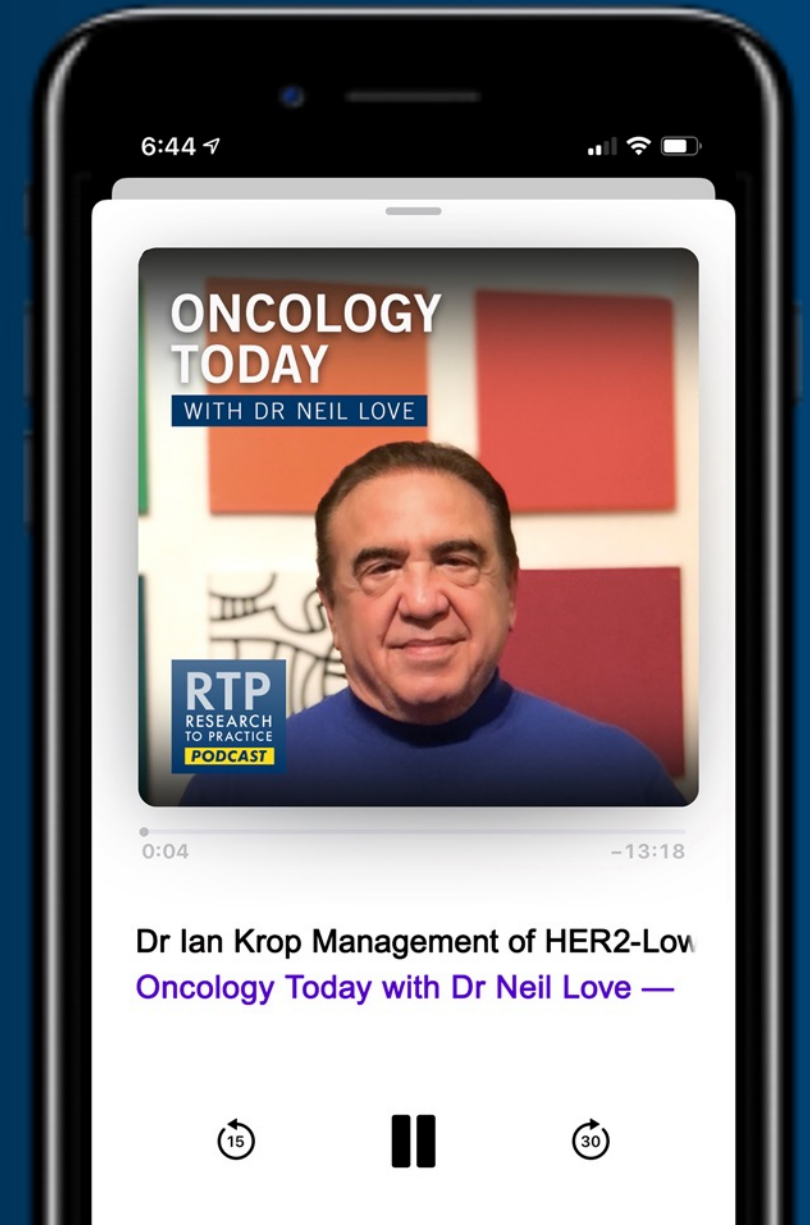
ONCOLOGY TODAY

WITH DR NEIL LOVE

Management of HER2-Low Breast Cancer



DR IAN KROP
DANA-FARBER CANCER INSTITUTE



Key Considerations in the Optimal Clinical Care of Patients with Small Cell Lung Cancer

A CME/MOC-Accredited Virtual Event

Thursday, November 4, 2021

5:00 PM – 6:00 PM ET

Faculty

Anne Chiang, MD, PhD

David R Spigel, MD

Moderator

Neil Love, MD

Meet The Professor

Optimizing the Management of Acute Myeloid Leukemia

Monday, November 8, 2021

5:00 PM – 6:00 PM ET

Faculty

Keith W Pratz, MD

Moderator

Neil Love, MD

Meet The Professor

Optimizing the Management of Metastatic Castration-Resistant Prostate Cancer

**Tuesday, November 9, 2021
5:00 PM – 6:00 PM ET**

Faculty

Simon Chowdhury, MD, PhD

Moderator

Neil Love, MD

VIRTUAL MOLECULAR TUMOR BOARD
Optimizing Biomarker-Based Decision-Making for
Patients with Non-Small Cell Lung Cancer with EGFR
Mutations or with Other Oncogene-Addicted Lung Cancers

A 2-Part CME/MOC-Accredited Webinar Series

Thursday, November 11, 2021

5:00 PM – 6:00 PM ET

Faculty

Marc Ladanyi, MD

Andrew J McKenzie, PhD

Helena Yu, MD

Moderator

Neil Love, MD

Meet The Professor
**Optimizing the Clinical Management of
Hodgkin and Non-Hodgkin Lymphomas**

**Monday, November 15, 2021
5:00 PM – 6:00 PM ET**

Faculty

Christopher R Flowers, MD, MS

Moderator

Neil Love, MD

Meet The Professor

Optimizing the Selection and Sequencing of Therapy for Patients with ER-Positive Breast Cancer

Wednesday, November 17, 2021

5:00 PM – 6:00 PM ET

Faculty

Kevin Kalinsky, MD, MS

Moderator

Neil Love, MD

Thank you for joining us!

CME and MOC credit information will be emailed to each participant within 5 business days.

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Meet The Professor Program Participating Faculty



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We Encourage Clinicians in Practice to Submit Questions

The screenshot displays a Zoom meeting interface. At the top, a gallery view shows six participants. The main area features a presentation slide with the text: "You may submit questions using the Zoom Chat option below" and a large red arrow pointing downwards. To the right, a "Participants (10)" list is visible, showing names like John Smith, Mary Major, Richard Miles, John Noakes, and Alice Suarez. A "Zoom Group Chat" window is open in the foreground, showing a message from "Me to Everyone" at 12:49 PM. The bottom toolbar includes icons for "Join Audio", "Start Video", "Invite", "Participants", "Share", "Chat", "Record", "Leave Meeting", "Mute Me", and "Raise Hand".

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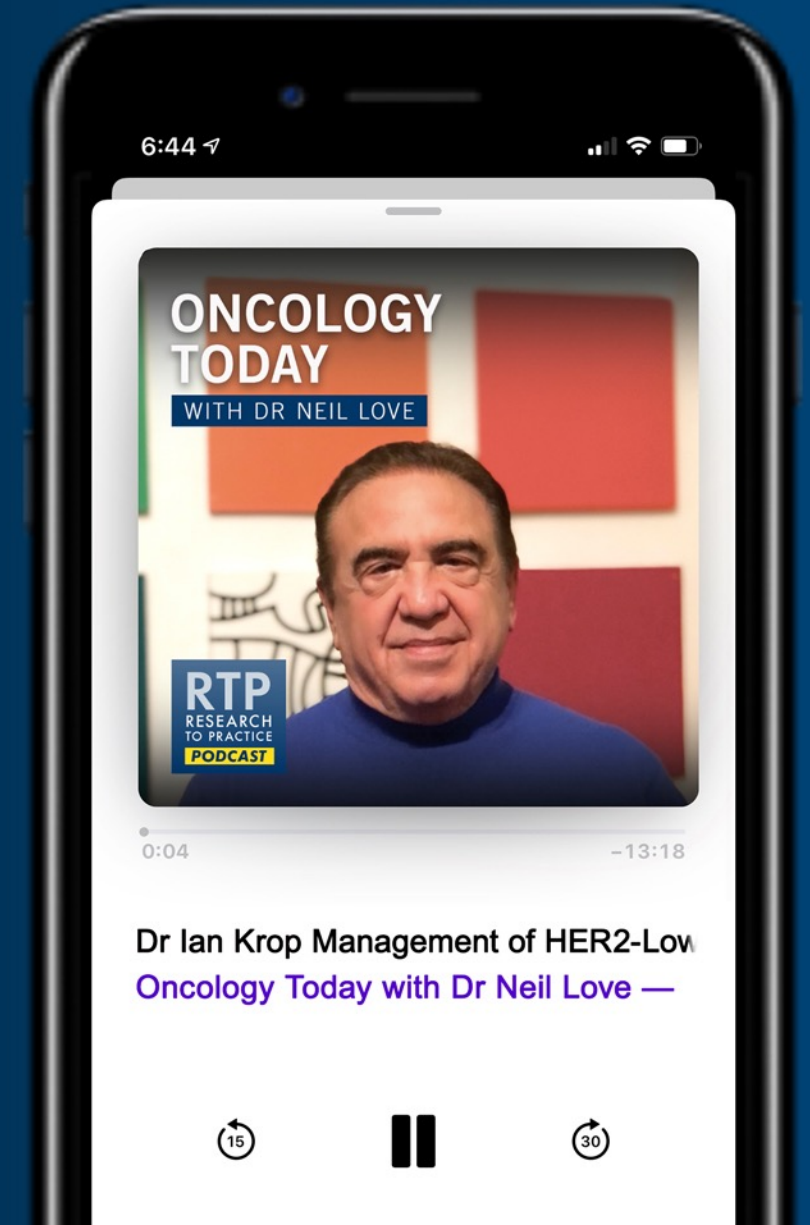
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A 65-year-old woman with an ER-positive, HER2-positive IDC experiences recurrence in the liver and brain 18 months after completing neoadjuvant TCHP followed by adjuvant trastuzumab/pertuzumab and postadjuvant neratinib and is receiving adjuvant anastrozole. Regulatory and reimbursement issues aside, what systemic treatment would you recommend?



Dr Gelmon

**Tucatinib +
trastuzumab/
capecitabine**



Dr Mahtani

**Trastuzumab
deruxtecan**



Dr Hamilton

**Tucatinib +
trastuzumab/
capecitabine**



Dr Rugo

**Trastuzumab
deruxtecan**



Dr Hurvitz

**Tucatinib +
trastuzumab/
capecitabine**



Dr Tolaney

**Trastuzumab/
pertuzumab/paclitaxel**



Alan B Astrow, MD
NewYork-Presbyterian Brooklyn
Methodist Hospital
Weill Cornell Medical College
Brooklyn, New York



Shachar Peles, MD
Florida Cancer Specialists
and Research Institute
Lake Worth, Florida



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Joseph Martins, MD
UT Health Science Center
Tyler, Texas

Meet The Professor with Dr Brufsky

MODULE 1: HER2 in the Real World

MODULE 2: Case Presentations

- Dr Martins: A 60-year-old woman with triple-positive metastatic breast cancer (mBC)
- Dr Gosain: A 67-year-old woman with ER/PR-negative, HER2-positive BC and brain metastases
- Dr Astrow: A 67-year-old woman with ER/PR-positive, HER2-positive mBC with recurrence in the brain
- Dr Peles: An 80-year-old woman with ER-positive, PR-negative, HER2-positive localized BC and treated CLL
- Dr Gandhi: A 58-year-old woman with ER/PR-negative, HER2-positive mBC with a PIK3CA mutation
- Dr Santiago: A 53-year-old woman with ER/PR-positive, HER2-positive infiltrating lobular BC
- Dr Zafar: A 33-year-old woman with triple-positive mBC and possible mosaicism of TP53 mutation

MODULE 3: Journal Club with Dr Brufsky

MODULE 4: Faculty Survey

MODULE 5: Appendix of Key Data Sets

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Breast Cancer (Dove Med Press) 2021;13:199-211

Breast Cancer: Targets and Therapy

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REVIEW

Approaching Neoadjuvant Therapy in the Management of Early-Stage Breast Cancer

Tara Hyder¹

Saveri Bhattacharya^{2,*}

Kristine Gade^{3,*}

Azadeh Nasrazadani³


Adam M Brufsky³

Breast Cancer Research and Treatment (2021) 188:179–190

<https://doi.org/10.1007/s10549-021-06103-z>

EPIDEMIOLOGY

Baseline characteristics and first-line treatment patterns in patients with HER2-positive metastatic breast cancer in the SystHERs registry

Peter A. Kaufman¹  · Sara A. Hurvitz² · Joyce O'Shaughnessy³ · Ginny Mason⁴ · Denise A. Yardley⁵ · Adam M. Brufsky⁶ · Hope S. Rugo⁷ · Melody Cobleigh⁸ · Sandra M. Swain⁹ · Debu Tripathy¹⁰ · Anne Morris¹¹ · Vincent Antao¹¹ · Haocheng Li¹² · Mohammad Jahanzeb¹³

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Case Presentation – Dr Martins: A 60-year-old woman with triple-positive mBC



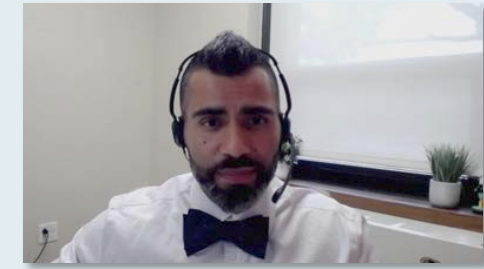
Dr Joseph Martins

- 2014: Triple-positive breast cancer with asymptomatic osseous metastases
- Taxane/trastuzumab/pertuzumab → Tamoxifen/trastuzumab
 - Ejection fraction dropped to 45% but normalized on beta-B and ACE-I
- 2/2016: Anastrozole
- 4/2016: PD → Resumed trastuzumab/pertuzumab
- 2/2019: PD → T-DM1, discontinued anastrozole
- 4/2020: PD → Switched to trastuzumab deruxtecan

Question

- What treatment would you consider next for her?

Case Presentation – Dr Gosain: A 67-year-old woman with ER/PR-negative, HER2-positive breast cancer and brain metastases



Dr Rohit Gosain

- Presented with 20-pound weight loss in 3 months and concerning mammogram finding in right breast
- Ultrasound and biopsy revealed a 5-cm IDC, HER2+ (positive based on FISH), ER/PR-negative
- CT of the chest/abdomen/pelvis and bone scan showed multiple liver and lung lesions
- THP (docetaxel/trastuzumab/pertuzumab) followed by T-DM1
- Presented with altered mental status and MRI showing multiple sub-centimeter brain lesions concerning for metastases, along with liver and lung lesions increased in size

Question

- What therapy would you offer next to this patient?

Case Presentation – Dr Astrow: A 67-year-old woman with ER/PR-positive, HER2-positive mBC with recurrence in the brain



Dr Alan Astrow

- Presented 6 years ago with triple-positive metastatic breast cancer and liver failure from liver metastases
- Carboplatin plus trastuzumab and an aromatase inhibitor; paclitaxel added as liver function improved
- Carboplatin/paclitaxel/trastuzumab x 6 months → continued aromatase inhibitor + trastuzumab/pertuzumab → CR for 5 years
- Developed brain metastases and treated with SBRT twice
- Now presents with new brain metastases

Questions

- Would you recommend another treatment with stereotactic brain radiation, or tucatinib plus capecitabine?
- What is your experience with tucatinib? How toxic is it? How are the patients doing on it? How long do you need to keep the patient on it? Would you combine it with SBRT?

Case Presentation – Dr Peles: An 80-year-old woman with ER-positive, PR-negative, HER2-positive localized breast cancer and treated CLL



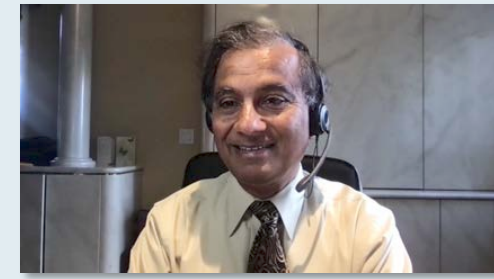
Dr Shachar Peles

- PMH: CLL, Rai Stage IV, s/p obinutuzumab/venetoclax 3/2021, CAD, CABG 2018; Recent EF: 65%
- Right, 2.7-cm ER-positive, PR-negative, HER2-positive IDC
- Discussed neoadjuvant paclitaxel/trastuzumab → Surgery → Trastuzumab x 1 year

Questions

- How would you manage her if she were 30 years younger? What if her ejection fraction was 40%?

Case Presentation – Dr Gandhi: A 58-year-old woman with ER/PR-negative, HER2-positive mBC with a PIK3CA mutation



Dr Sunil Gandhi

- 2-cm ER-negative, PR-positive, HER2-positive BC s/p lumpectomy
- Paclitaxel/trastuzumab x 12 → Trastuzumab x 1 year; Declined adjuvant anastrozole
- Transferred care 6/2021: Hepatomegaly, with widespread biopsy-confirmed ER/PR-negative, HER2-positive liver metastases
- Docetaxel/carboplatin/trastuzumab/pertuzumab (TCHP), with near CR after 3 cycles

Question

- When should we administer neoadjuvant chemotherapy versus adjuvant chemotherapy for patients with HER2-positive breast cancer?

Case Presentation – Dr Santiago: A 53-year-old woman with ER/PR-positive, HER2-positive infiltrating lobular breast cancer



Dr Ferdy Santiago

- Currently receiving neoadjuvant TCHP for a 2.5-cm, Grade II, ER/PR/HER2-positive infiltrating lobular cancer

Question

- If she achieves a pathologic complete response but has circulating tumor cells, would you administer T-DM1 as per the KATHERINE trial?

Case Presentation – Dr Zafar: A 33-year-old woman with triple-positive mBC and possible mosaicism of TP53 mutation



Dr Syed Zafar

- Bilateral breast masses and LAD
 - Right: 6-cm, Grade III, ER/PR/HER2-positive
 - Left: 3-cm, Grade III, ER/PR-positive, HER2-negative
- Solitary 1.7-cm right hepatic metastasis, biopsy-confirmed ER/PR/HER2-positive
- Germline testing: BRCA wildtype, possible mosaicism of TP53 mutation
- OFS + trastuzumab/pertuzumab/docetaxel, with good response (6-mm residual liver lesion, SBRT)
- Maintenance trastuzumab/pertuzumab + OFS/AI and bisphosphonate

Question

- Should we consider breast surgery of her primaries combined with local therapy, or just proceed with her current maintenance treatment?

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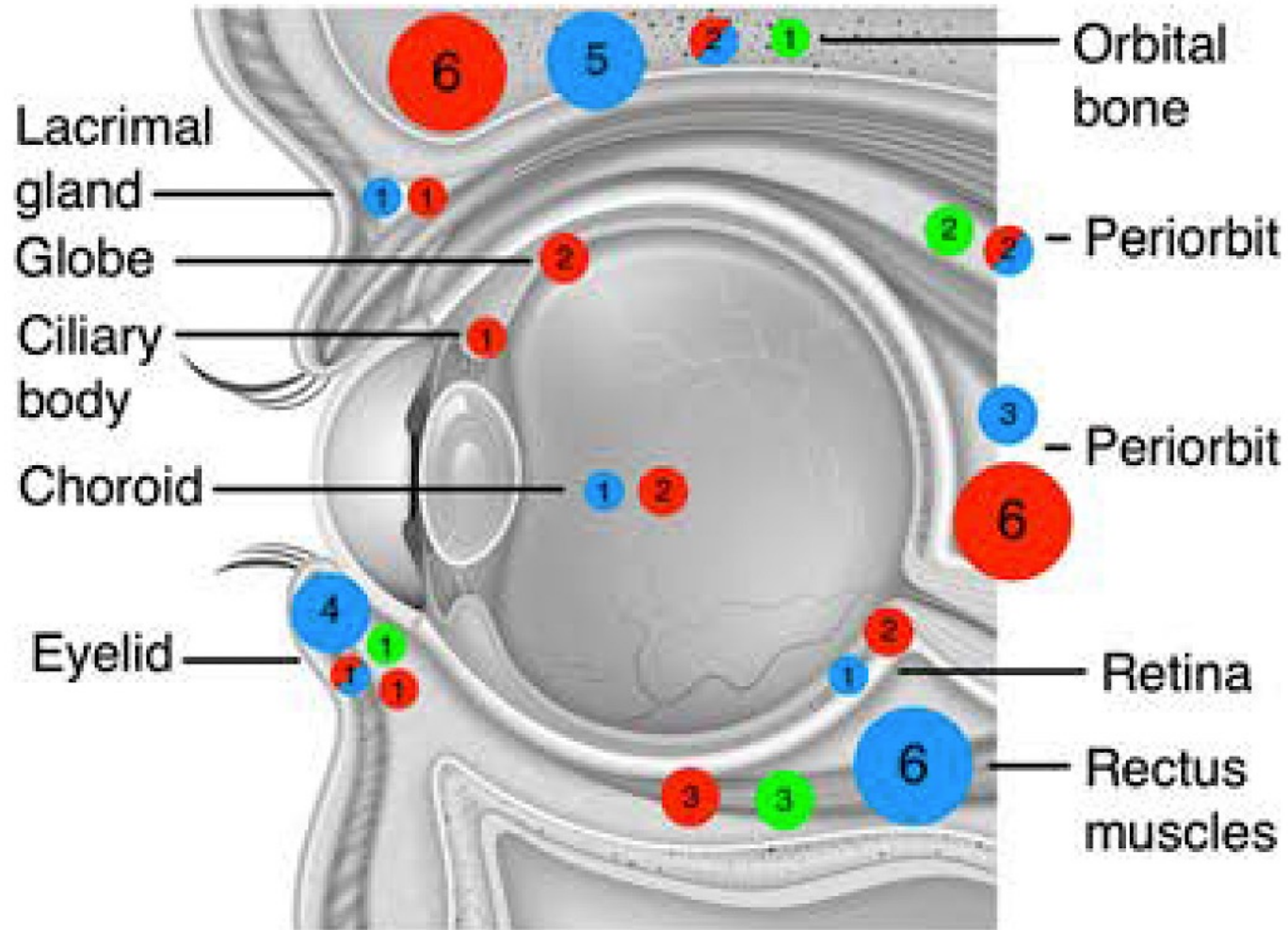
RESEARCH ARTICLE

Open Access

Patient treatment and outcome after breast cancer orbital and periorbital metastases: a comprehensive case series including analysis of lobular versus ductal tumor histology

Martin Blohmer^{1,2,3}, Li Zhu⁴, Jennifer M. Atkinson^{1,3}, Sushil Beriwal⁵, Joshua L. Rodríguez-López⁵, Margaret Rosenzweig^{3,6}, Adam M. Brufsky⁷, George Tseng^{3,7}, Peter C. Lucas^{3,8}, Adrian V. Lee^{1,3†}, Steffi Oesterreich^{1,3†} and Rachel C. Jankowitz^{7,9,10*†}

Representation of the Anatomical Location of All Ophthalmic Metastases (OM)



Red dots represent metastases from an IDC primary, blue dots represent metastases from an ILC primary, mixed red and blue dots represent metastases from a mixed IDC/ILC primary, and green dots represent metastases from a primary of unknown histological subtype.

Numbers indicate how many patients were affected by OM to this location. In cases where patients had OM to multiple locations within the ophthalmic region, each location was displayed separately.

Journal Club with Dr Brufsky

- Brufsky A et al. **A Phase 2 study of poziotinib in patients with HER2-positive metastatic breast cancer heavily pre-treated with HER2-targeted therapy.** SABCS 2020;Abstract PD1-07.
- Hurvitz SA et al. **Efficacy of neratinib plus capecitabine in the subgroup of patients with central nervous system involvement from the NALA trial.** *Oncologist* 2021;26(8):e1327-38.
- Marx GM et al. **Dose escalation for mitigating diarrhea: Ranked tolerability assessment of anti-diarrheal regimens in patients receiving neratinib for early-stage breast cancer.** ASCO 2021;Abstract 536.
- Hyder T et al. **Aromatase inhibitor-associated musculoskeletal syndrome: Understanding mechanisms and management.** *Front Endocrinol (Lausanne)* 2021;12:713700.

Journal Club with Dr Brufsky

- Mamounas EP et al. **Breast Cancer Index (BCI) and prediction of benefit from extended aromatase inhibitor (AI) therapy (tx) in HR+ breast cancer: NRG oncology/NSABP B-42.** ASCO 2021;Abstract 501.
- Lundstrom K et al. **Viewpoint: Origin of SARS-CoV-2.** *Viruses* 2020;12(11):1203.
- Redwan EM et al. **The mechanism behind flaring/triggering of autoimmunity disorders associated with COVID-19.** *Autoimmun Rev* 2021;20(10):102909.
- Perrot L et al. **First flare of ACPA-positive rheumatoid arthritis after SARS-CoV-2 infection.** *Lancet Rheumatol* 2021;3(1):e6-8.

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Management of Metastatic HER2-Positive Breast Cancer

A 65-year-old woman with an ER-negative, HER2-positive IDC experiences disease recurrence in the liver 6 months after completing neoadjuvant TCHP followed by adjuvant trastuzumab/pertuzumab. Regulatory and reimbursement issues aside, what systemic treatment would you recommend?

1. Trastuzumab/pertuzumab/docetaxel
2. T-DM1
3. Neratinib + paclitaxel
4. Neratinib + capecitabine
5. Tucatinib + trastuzumab/capecitabine
6. Trastuzumab deruxtecan
7. Trastuzumab + capecitabine
8. Other

A 65-year-old woman with an ER-negative, HER2-positive IDC experiences disease recurrence in the liver 6 months after completing neoadjuvant TCHP followed by adjuvant trastuzumab/pertuzumab. Regulatory and reimbursement issues aside, what systemic treatment would you recommend?



Dr Gelmon

**Trastuzumab
deruxtecan**



Dr Mahtani

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Dr Hamilton

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Dr Rugo

**Trastuzumab
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Dr Hurvitz

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deruxtecan**



Dr Tolaney

**Trastuzumab
deruxtecan**

A 65-year-old woman with an ER-negative, HER2-positive IDC experiences disease recurrence in the liver 6 months after completing neoadjuvant TCHP followed by adjuvant T-DM1. Regulatory and reimbursement issues aside, what systemic treatment would you recommend?

1. Trastuzumab/pertuzumab/docetaxel
2. Neratinib + paclitaxel
3. Neratinib + capecitabine
4. Tucatinib + trastuzumab/capecitabine
5. Trastuzumab deruxtecan
6. Trastuzumab + capecitabine
7. Other

A 65-year-old woman with an ER-negative, HER2-positive IDC experiences disease recurrence in the liver 6 months after completing neoadjuvant TCHP followed by adjuvant T-DM1. Regulatory and reimbursement issues aside, what systemic treatment would you recommend?



Dr Gelmon

**Trastuzumab
deruxtecan**



Dr Mahtani

**Trastuzumab
deruxtecan**



Dr Hamilton

**Trastuzumab
deruxtecan**



Dr Rugo

**Trastuzumab
deruxtecan**



Dr Hurvitz

**Trastuzumab
deruxtecan**



Dr Tolaney

**Trastuzumab
deruxtecan**

A 65-year-old woman with an ER-negative, HER2-positive IDC experiences disease recurrence in the liver and brain 18 months after completing neoadjuvant TCHP followed by adjuvant trastuzumab/pertuzumab. Regulatory and reimbursement issues aside, what systemic treatment would you recommend?

1. Trastuzumab/pertuzumab/docetaxel
2. T-DM1
3. Neratinib + paclitaxel
4. Neratinib + capecitabine
5. Tucatinib + trastuzumab/capecitabine
6. Trastuzumab deruxtecan
7. Trastuzumab + capecitabine
8. Other

A 65-year-old woman with an ER-negative, HER2-positive IDC experiences disease recurrence in the liver and brain 18 months after completing neoadjuvant TCHP followed by adjuvant trastuzumab/pertuzumab. Regulatory and reimbursement issues aside, what systemic treatment would you recommend?



Dr Gelmon

**Tucatinib +
trastuzumab/
capecitabine**



Dr Mahtani

**Tucatinib +
trastuzumab/
capecitabine**



Dr Hamilton

**Tucatinib +
trastuzumab/
capecitabine**



Dr Rugo

**Tucatinib +
trastuzumab/
capecitabine**



Dr Hurvitz

**Trastuzumab
deruxtecan**



Dr Tolaney

**Trastuzumab/
pertuzumab/paclitaxel**

A 65-year-old woman with an ER-positive, HER2-positive IDC experiences recurrence in the liver and brain 18 months after completing neoadjuvant TCHP followed by adjuvant trastuzumab/pertuzumab and postadjuvant neratinib and is receiving adjuvant anastrozole. Regulatory and reimbursement issues aside, what systemic treatment would you recommend?



Dr Gelmon

**Tucatinib +
trastuzumab/
capecitabine**



Dr Mahtani

**Trastuzumab
deruxtecan**



Dr Hamilton

**Tucatinib +
trastuzumab/
capecitabine**



Dr Rugo

**Trastuzumab
deruxtecan**



Dr Hurvitz

**Tucatinib +
trastuzumab/
capecitabine**



Dr Tolaney

**Trastuzumab/
pertuzumab/
paclitaxel**

A 65-year-old woman with ER-negative, HER2-positive mBC receives first-line THP followed by second-line T-DM1 on disease progression. She now presents with further low-volume, asymptomatic progression but no evidence of CNS involvement. Regulatory and reimbursement issues aside, what systemic treatment would you recommend?



Dr Gelmon

**Tucatinib +
trastuzumab/
capecitabine**



Dr Mahtani

**Tucatinib +
trastuzumab/
capecitabine**



Dr Hamilton

**Tucatinib +
trastuzumab/
capecitabine**



Dr Rugo

**Trastuzumab
deruxtecan**



Dr Hurvitz

**Tucatinib +
trastuzumab/
capecitabine**



Dr Tolaney

**Trastuzumab
deruxtecan**

A 65-year-old woman with ER-negative, HER2-positive mBC receives first-line THP followed by second-line T-DM1 on disease progression. She now presents with further high-volume, moderately symptomatic progression but no evidence of CNS involvement. Regulatory and reimbursement issues aside, what systemic treatment would you recommend?



Dr Gelmon

**Tucatinib +
trastuzumab/
capecitabine**



Dr Mahtani

**Trastuzumab
deruxtecan**



Dr Hamilton

**Trastuzumab
deruxtecan**



Dr Rugo

**Trastuzumab
deruxtecan**



Dr Hurvitz






**Trastuzumab
deruxtecan**



Dr Tolaney

**Trastuzumab
deruxtecan**

At what grade of ILD would you permanently discontinue therapy with trastuzumab deruxtecan for a patient with HER2-positive mBC?

 Dr Gelmon	Grade 2	 Dr Mahtani	Grade 2
 Dr Hamilton	Grade 2	 Dr Rugo	Grade 2
 Dr Hurvitz	Grade 2	 Dr Tolaney	Grade 2

A 65-year-old woman with ER-negative, HER2-positive mBC receives first-line THP but after 1 year experiences disease progression, including 1 brain metastasis that is resected. Regulatory and reimbursement issues aside, what systemic treatment would you recommend next?



Dr Gelmon

Tucatinib +
trastuzumab/
capecitabine



Dr Mahtani

Tucatinib +
trastuzumab/
capecitabine



Dr Hamilton

Tucatinib +
trastuzumab/
capecitabine



Dr Rugo

Trastuzumab
deruxtecan



Dr Hurvitz

Trastuzumab
deruxtecan



Dr Tolaney

Trastuzumab
deruxtecan

A 65-year-old woman with ER-negative, HER2-positive mBC receives first-line THP but after 1 year experiences disease progression, including multiple brain metastases. Regulatory and reimbursement issues aside, what systemic treatment would you recommend next?



Dr Gelmon

**Tucatinib +
trastuzumab/
capecitabine**



Dr Mahtani

**Tucatinib +
trastuzumab/
capecitabine**



Dr Hamilton

**Tucatinib +
trastuzumab/
capecitabine**



Dr Rugo

**Tucatinib +
trastuzumab/
capecitabine**



Dr Hurvitz

**Tucatinib +
trastuzumab/
capecitabine**



Dr Tolaney

**Tucatinib +
trastuzumab/
capecitabine**

A 65-year-old woman with ER-negative, HER2-positive mBC receives first-line THP followed by second-line T-DM1 on disease progression. She now presents with a single brain metastasis that is resected with no other evidence of progression. Regulatory and reimbursement issues aside, what systemic treatment would you recommend?



Dr Gelmon

Continue T-DM1



Dr Mahtani

Continue T-DM1



Dr Hamilton

Continue T-DM1



Dr Rugo

Continue T-DM1



Dr Hurvitz

Continue T-DM1



Dr Tolaney

Continue T-DM1

A 65-year-old woman with ER-negative, HER2-positive mBC receives first-line THP followed by second-line T-DM1 on disease progression. She now presents with further disease progression, including multiple new brain metastases. Regulatory and reimbursement issues aside, what systemic treatment would you recommend?



Dr Gelmon

**Tucatinib +
trastuzumab/
capecitabine**



Dr Hamilton

**Tucatinib +
trastuzumab/
capecitabine**



Dr Hurvitz

**Tucatinib +
trastuzumab/
capecitabine**



Dr Mahtani

**Tucatinib +
trastuzumab/
capecitabine**



Dr Rugo

**Tucatinib +
trastuzumab/
capecitabine**



Dr Tolaney

**Tucatinib +
trastuzumab/
capecitabine**

Localized HER2-Positive Breast Cancer

Which neoadjuvant systemic therapy, if any, would you generally recommend for a 65-year-old patient with a 2.5-cm ER-negative, HER2-positive, clinically node-negative IDC?

1. None

2. TCHP

3. TCH







4. Paclitaxel/trastuzumab

5. Paclitaxel/trastuzumab/pertuzumab

6. ACTH

7. Other

Which neoadjuvant systemic therapy, if any, would you generally recommend for a 65-year-old patient with a 2.5-cm ER-negative, HER2-positive, clinically node-negative IDC?

 Dr Gelmon	TCHP (TCH/pertuzumab) or ACTH/pertuzumab	 Dr Mahtani	TCHP
 Dr Hamilton	TCHP	 Dr Rugo	Paclitaxel/trastuzumab /pertuzumab
 Dr Hurvitz	TCHP	 Dr Tolaney	TCHP

A 65-year-old woman presents with a 3.4-cm ER-positive, HER2-positive IDC with biopsy-proven axillary nodes, receives neoadjuvant TCHP and at surgery is found to have 0.5 cm of residual tumor in the breast and no disease in the nodes. Regulatory and reimbursement issues aside, what adjuvant anti-HER2 therapy would you recommend?

1. Trastuzumab
2. Trastuzumab/pertuzumab
3. T-DM1
4. Trastuzumab → neratinib
5. Trastuzumab/pertuzumab → neratinib
6. T-DM1 → neratinib
7. Other

A 65-year-old woman presents with a 3.4-cm ER-positive, HER2-positive IDC with biopsy-proven axillary nodes, receives neoadjuvant TCHP and at surgery is found to have 0.5 cm of residual tumor in the breast and no disease in the nodes. Regulatory and reimbursement issues aside, what adjuvant anti-HER2 therapy would you recommend?



Dr Gelmon

T-DM1



Dr Mahtani

T-DM1



Dr Hamilton

**T-DM1 or
T-DM1 → neratinib**



Dr Rugo

T-DM1



Dr Hurvitz

T-DM1 → neratinib



Dr Tolaney

T-DM1

Meet The Professor with Dr Brufsky

MODULE 1: HER2 in the Real World

MODULE 2: Case Presentations

- Dr Martins: A 60-year-old woman with triple-positive metastatic breast cancer (mBC)
- Dr Gosain: A 67-year-old woman with ER/PR-negative, HER2-positive BC and brain metastases
- Dr Astrow: A 67-year-old woman with ER/PR-positive, HER2-positive mBC with recurrence in the brain
- Dr Peles: An 80-year-old woman with ER-positive, PR-negative, HER2-positive localized BC and treated CLL
- Dr Gandhi: A 58-year-old woman with ER/PR-negative, HER2-positive mBC with a PIK3CA mutation
- Dr Santiago: A 53-year-old woman with ER/PR-positive, HER2-positive infiltrating lobular BC
- Dr Zafar: A 33-year-old woman with triple-positive mBC and possible mosaicism of TP53 mutation

MODULE 3: Journal Club with Dr Brufsky

MODULE 4: Faculty Survey

MODULE 5: Appendix of Key Data Sets

Management of Metastatic HER2-Positive Breast Cancer

Trastuzumab Deruxtecan Significantly Improved PFS Over T-DM1 for HER2-Positive Metastatic Breast Cancer

Press Release – August 9, 2021

“Trastuzumab deruxtecan demonstrated superior progression-free survival (PFS) outcomes over trastuzumab emtansine (T-DM1) in patients with HER2-positive metastatic breast cancer, based on the phase 3 DESTINY-Breast03 trial (NCT03529110). The study’s planned interim analysis identified a statistically significant and clinically meaningful improvement in the primary end point of PFS as assessed by an Independent Data Monitoring Committee (IDMC) for patients with HER2-positive, unresectable and/or metastatic breast cancer who received prior treatment with trastuzumab and a taxane.

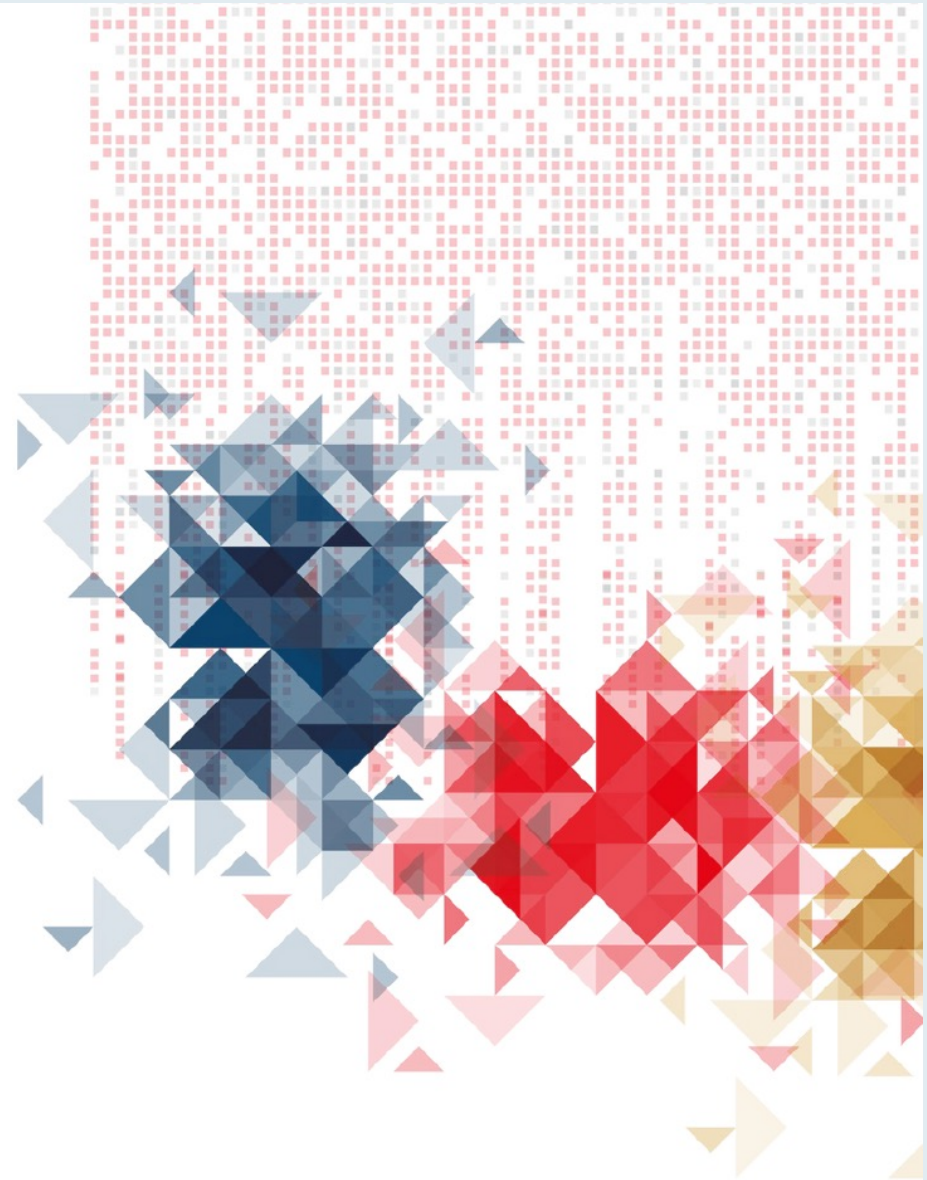
Approximately 500 patients were enrolled in the DESTINY-Breast03 trial, who were randomized to either the experimental trastuzumab deruxtecan arm or the comparator T-DM1 arm. The primary end point was PFS assessed by IDMC, with secondary end points including overall survival (OS), objective response rate (ORR), duration of response, and PFS based on investigator assessment.

While patients treated with trastuzumab deruxtecan trended toward OS improvement, the data were immature. Furthermore, the safety profile was consistent with previously reported data regarding trastuzumab deruxtecan, with no new safety signals or grade 4/5 treatment-related interstitial lung disease events observed.”

Trastuzumab Deruxtecan (T-DXd) vs Trastuzumab Emtansine (T-DM1) in Patients With HER2+ Metastatic Breast Cancer: Results of the Randomized, Phase 3 Study DESTINY-Breast03

Javier Cortés, MD^a, Sung-Bae Kim, Wei-Pang Chung, Seock-Ah Im, Yeon Hee Park, Roberto Hegg, Min-Hwan Kim, Ling-Ming Tseng, Vanessa Petry, Chi-Feng Chung, Hiroji Iwata, Erika Hamilton, Giuseppe Curigliano, Binghe Xu, Caleb Lee, Yali Liu, Jillian Cathcart, Emarjola Bako, Sunil Verma, Sara Hurvitz
On behalf of the DESTINY-Breast03 investigators

^aMedical Oncology, International Breast Cancer Center (IBCC), Quironsalud Group, and Vall d'Hebron Institute of Oncology (VHIO), Barcelona, Spain; Universidad Europea de Madrid, Faculty of Biomedical and Health Sciences, Department of Medicine, Madrid, Spain.



DESTINY-Breast03 Phase III Trial Schema

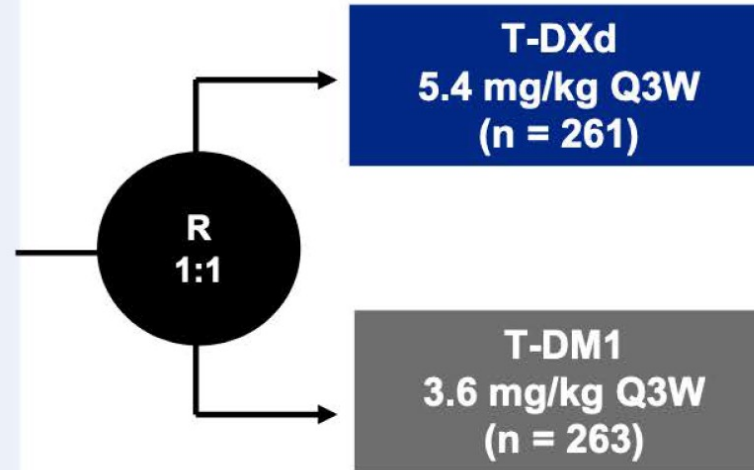
An open-label, multicenter study (NCT03529110)

Patients

- Unresectable or metastatic HER2-positive^a breast cancer
- Previously treated with trastuzumab and taxane in advanced/metastatic setting^b
- Could have clinically stable, treated brain metastases

Stratification factors

- Hormone receptor status
- Prior treatment with pertuzumab
- History of visceral disease



Primary endpoint

- PFS (BICR)

Key secondary endpoint

- OS

Secondary endpoints

- ORR (BICR and investigator)
- DOR (BICR)
- PFS (investigator)
- Safety

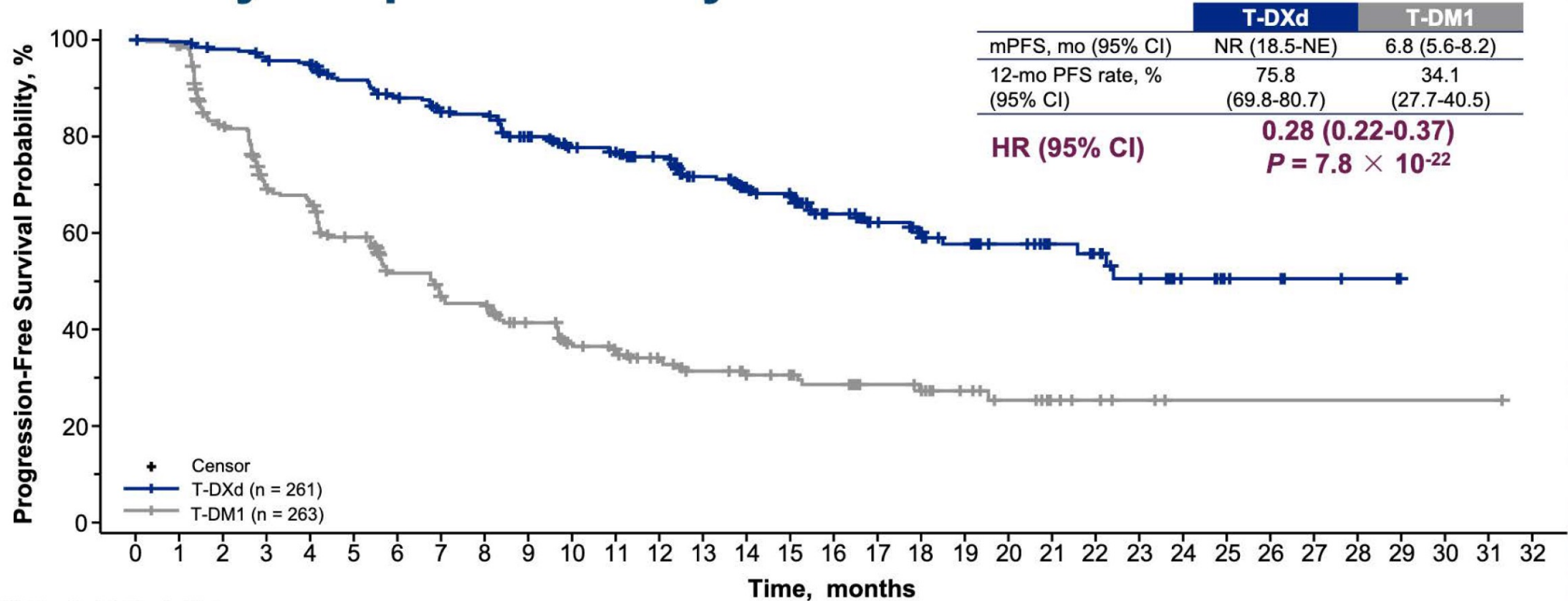
Interim analysis for PFS (data cutoff: May 21, 2021)

- Efficacy boundary for superiority: $P < 0.000204$ (based on 245 events)
- IDMC recommendation to unblind study (July 30, 2021)

Key secondary endpoint, OS: boundary for efficacy: $P < 0.000265$ (based on 86 events)

DESTINY-Breast03: PFS by BICR

Primary Endpoint: PFS by BICR

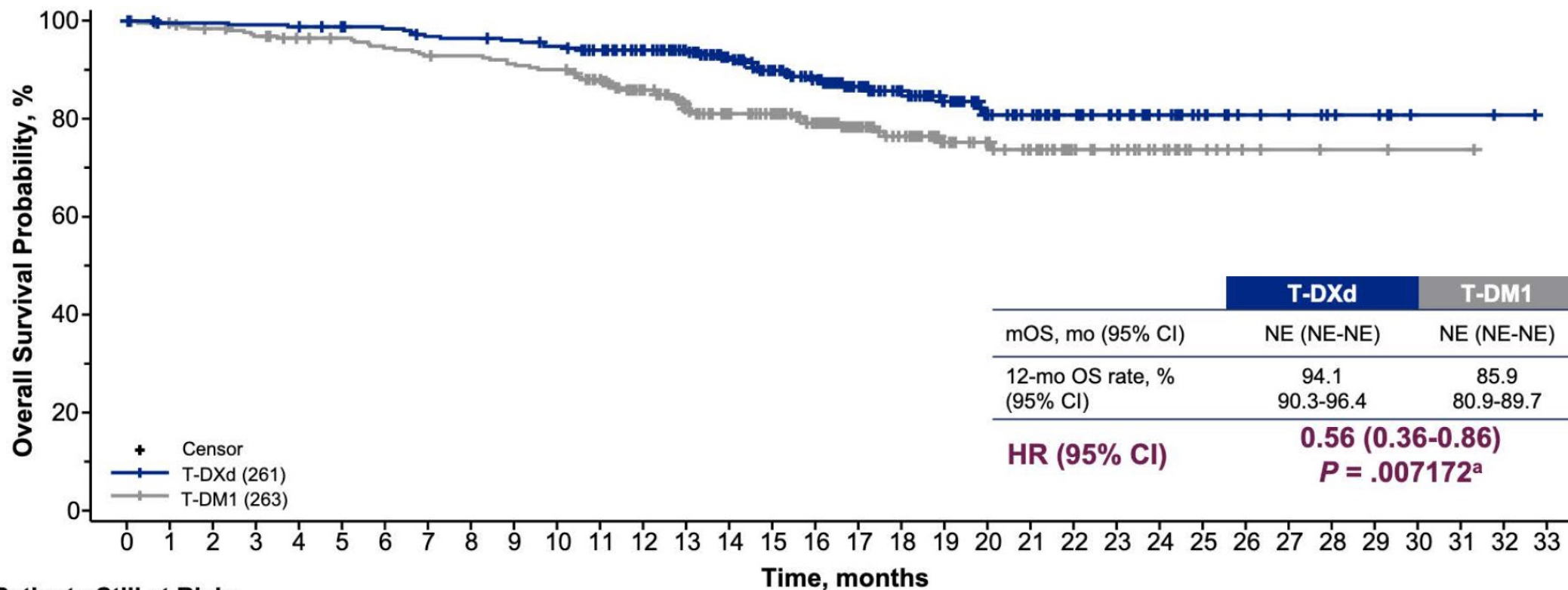


Patients Still at Risk:

T-DXd (261)	261	256	250	244	240	224	214	202	200	183	168	164	150	132	112	105	79	64	53	45	36	29	25	19	10	6	5	3	2	0			
T-DM1 (263)	263	252	200	163	155	132	108	96	93	78	65	60	51	43	37	34	29	23	21	16	12	8	6	4	1	1	1	1	1	1	1	1	0

DESTINY-Breast03: OS by BICR

Key Secondary Endpoint: OS



Patients Still at Risk:

T-DXd (261)	261	256	256	255	254	251	249	244	243	241	237	230	218	202	180	158	133	108	86	71	56	50	42	33	24	18	11	10	7	6	2	2	1	0
T-DM1 (263)	263	258	253	248	243	241	236	232	231	227	224	210	188	165	151	140	120	91	75	58	52	44	32	27	18	11	5	4	3	3	1	1	0	



Early OS data with relatively few events (33 in the T-DXd arm, 53 in the T-DM1 arm)

^aP = .007172, but does not cross pre-specified boundary of P < .000265



DESTINY-Breast03: Adverse Events of Special Interest

Adjudicated as drug-related ILD/pneumonitis ^a , n (%)						
n (%)	Grade 1	Grade 2	Grade 3	Grade 4	Grade 5	Any Grade
T-DXd (n = 257)	7 (2.7)	18 (7.0)	2 (0.8)	0	0	27 (10.5)
T-DM1 (n = 261)	4 (1.5)	1 (0.4)	0	0	0	5 (1.9)

- There were no grade 4 or 5 adjudicated drug-related ILD/pneumonitis events observed with T-DXd

LVEF decrease, n (%)						
n (%)	Grade 1	Grade 2	Grade 3	Grade 4	Grade 5	Any Grade
T-DXd (n = 257)	1 (0.4) ^b	6 (2.3) ^c	0	0	0	7 (2.7)
T-DM1 (n = 261)	0	1 (0.4) ^c	0	0	0	1 (0.4)

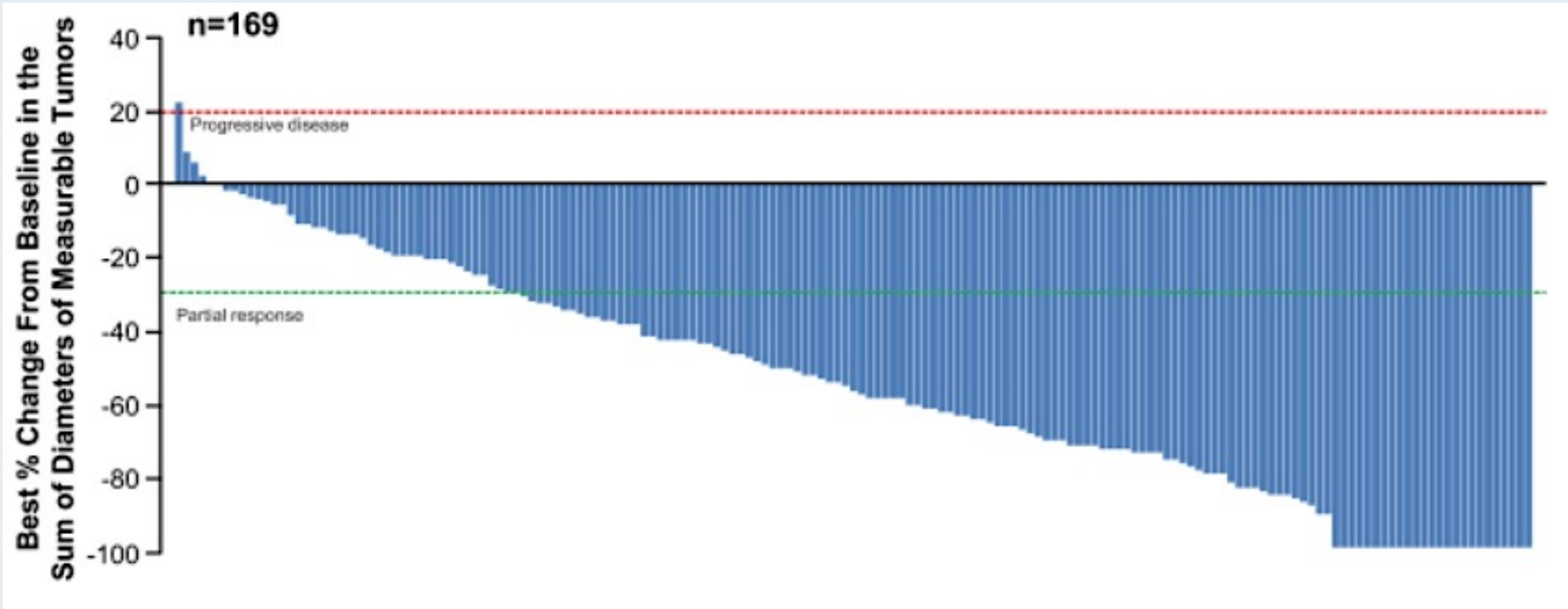
- In the T-DXd arm, all reported adverse events of LVEF decrease were asymptomatic and no cases of cardiac failure occurred

Updated Results from DESTINY-Breast01, a Phase 2 Trial of Trastuzumab Deruxtecan (T-DXd) in HER2-Positive Metastatic Breast Cancer

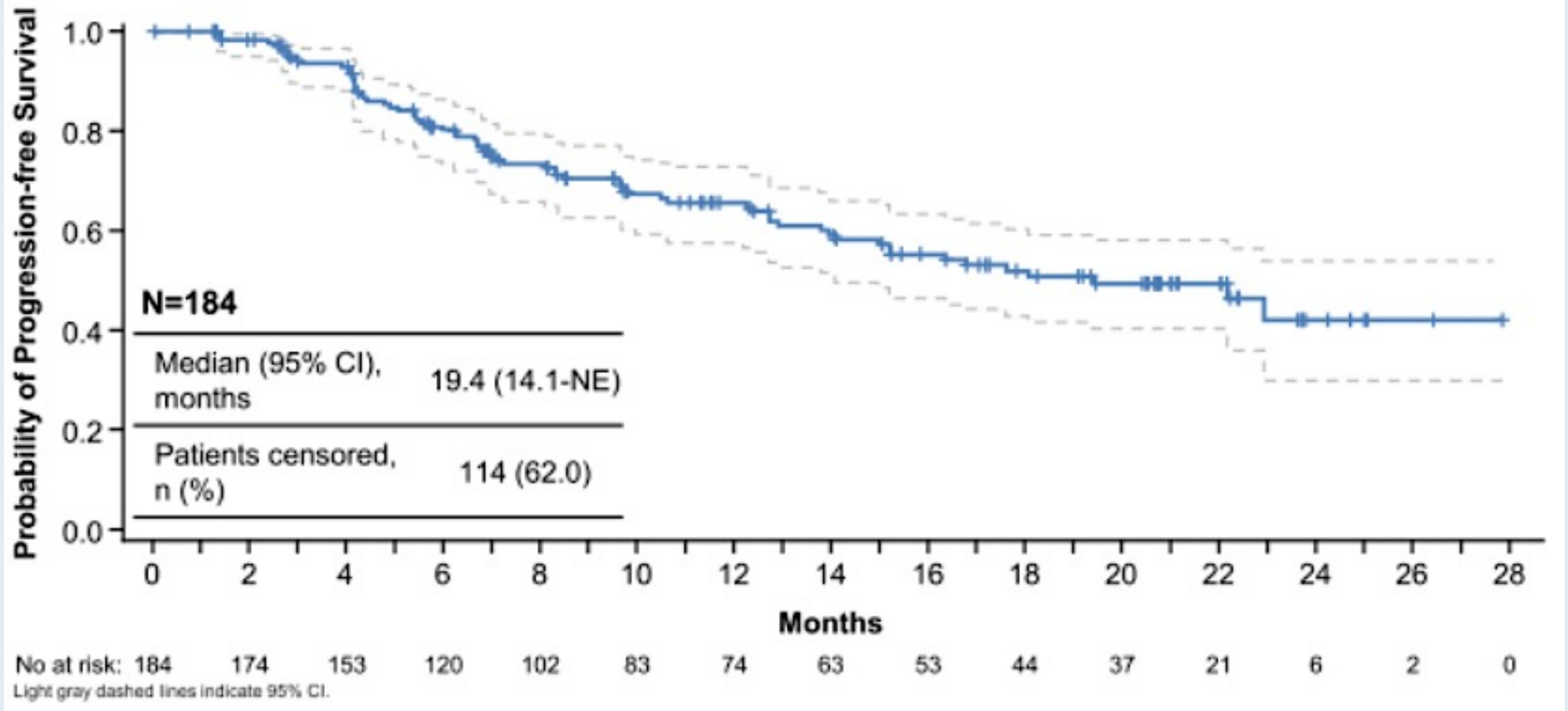
Modi S et al.

SABCS 2020;Abstract PD3-06.

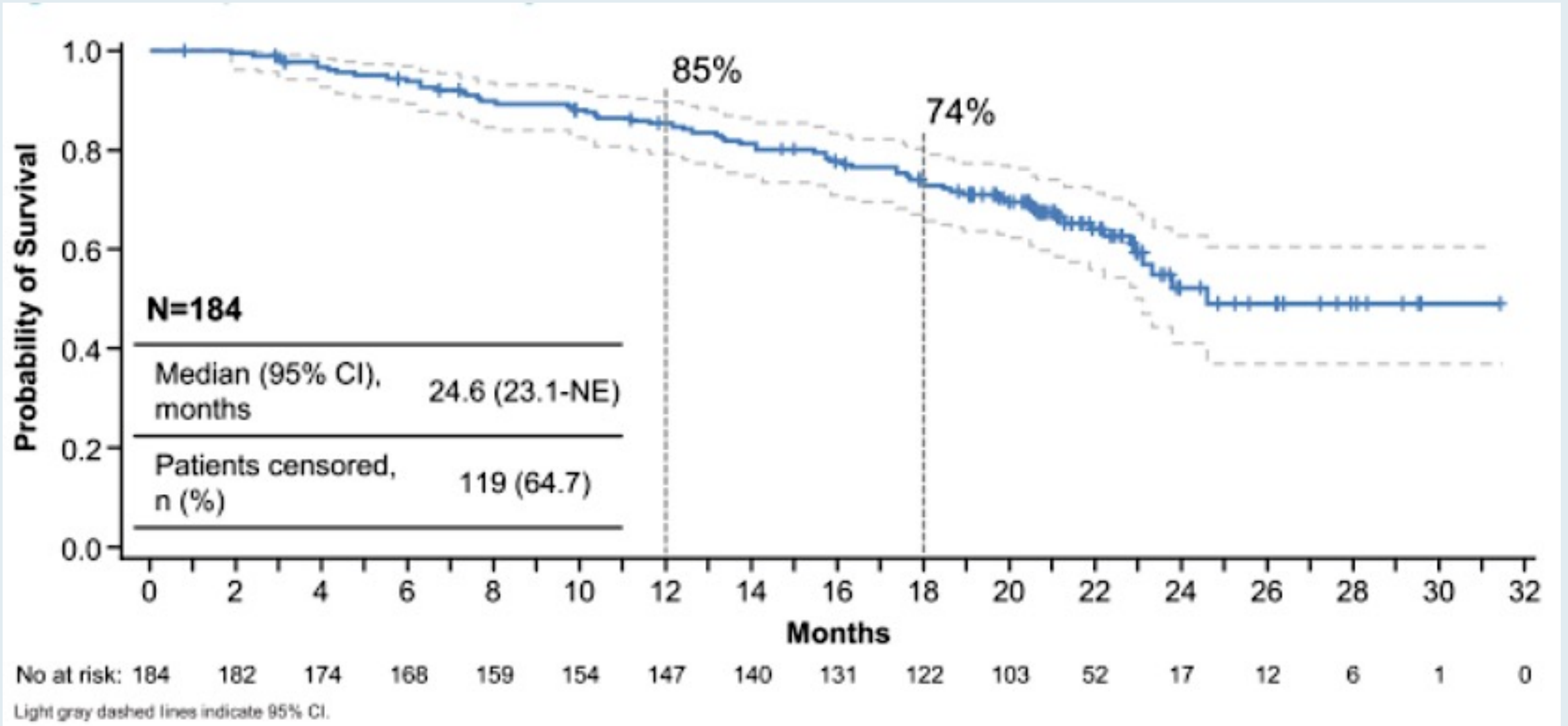
DESTINY-Breast01: Best Percent Change in Tumor Size from Baseline



DESTINY-Breast01: Progression-Free Survival



DESTINY-Breast01: Overall Survival



DESTINY-Breast01: Safety

AEs of special interest (n = 184)	All grades	Grades 3 and 4
Interstitial lung disease	25 (13.6%)	1 (0.5%)
Prolonged QT interval	9 (4.9%)	2 (1.1%)
Infusion-related reaction	4 (2.2%)	0
Decreased left ventricular ejection fraction	3 (1.6%)	1 (0.5%)

- Most common Grade ≥ 3 AEs were decreased neutrophil count (21%), anemia (9%) and nausea (8%).

Trastuzumab Deruxtecan (T-DXd) in Patients with HER2+ Metastatic Breast Cancer with Brain Metastases: A Subgroup Analysis of the DESTINY-Breast01 Trial

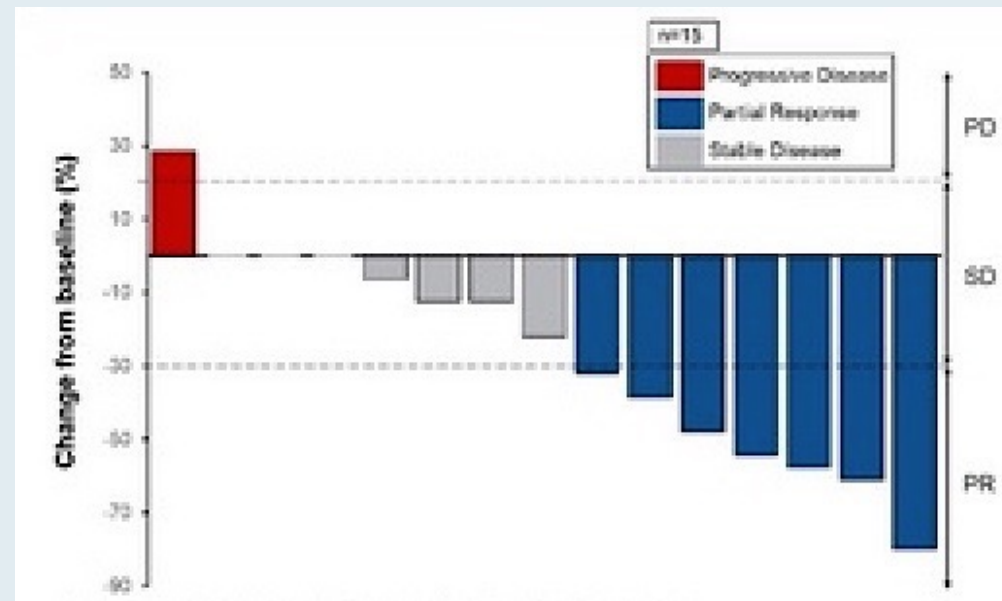
Jerusalem GHM et al.

ASCO 2021;Abstract 526.

DESTINY-Breast01: Clinical Activity Outcomes with Trastuzumab Deruxtecan

Endpoint	CNS Subgroup (n = 24)	All Patients (N = 184)
Confirmed ORR	58.3%	60.9%
Duration of response	16.9 mo	14.8 mo
Progression-free survival	18.1 mo	16.4 mo

Best Response in Brain Lesions in the CNS Subgroup

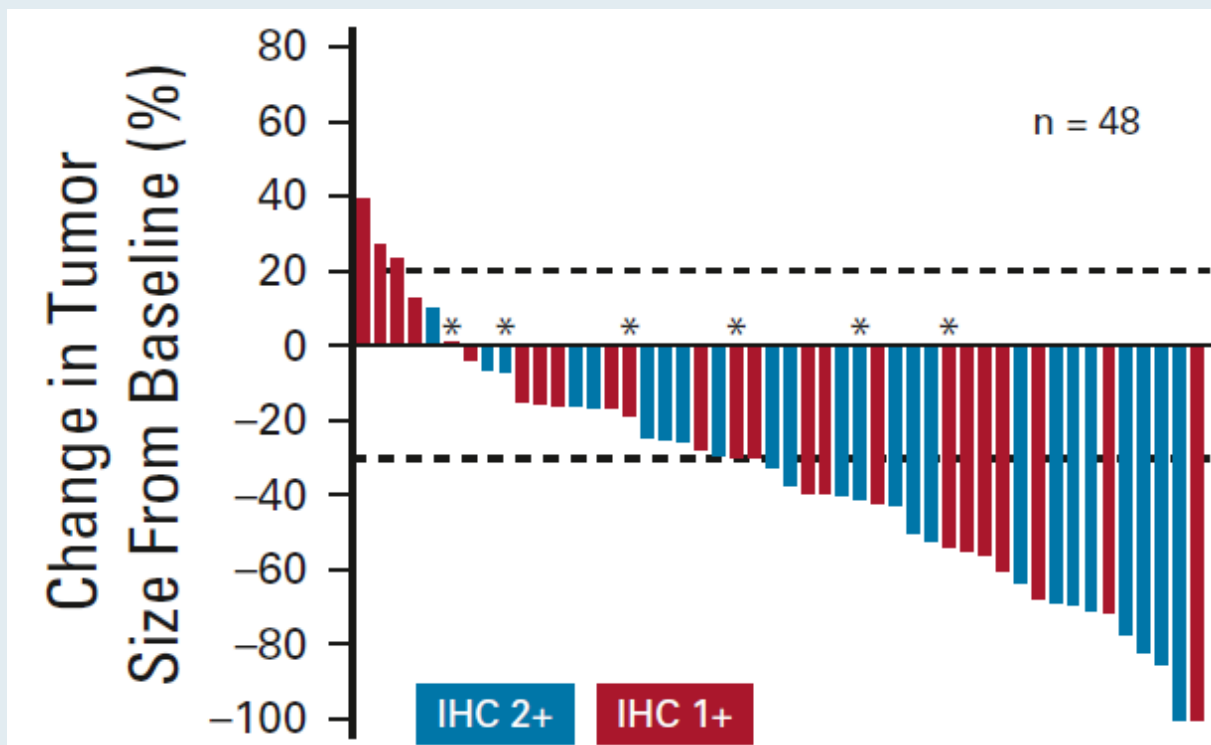


Antitumor Activity and Safety of Trastuzumab Deruxtecan in Patients With HER2-Low–Expressing Advanced Breast Cancer: Results From a Phase Ib Study

Shanu Modi, MD¹; Haeseong Park, MD, MPH²; Rashmi K. Murthy, MD, MBE³; Hiroji Iwata, PhD, MD⁴; Kenji Tamura, MD, PhD⁵; Junji Tsurutani, MD, PhD⁶; Alvaro Moreno-Aspitia, PhD⁷; Toshihiko Doi, MD, PhD⁸; Yasuaki Sagara, MD⁹; Charles Redfern, MD¹⁰; Ian E. Krop, MD, PhD¹¹; Caleb Lee, MD, PhD¹²; Yoshihiko Fujisaki, MS¹³; Masahiro Sugihara, PhD¹³; Lin Zhang, MD, PhD¹²; Javad Shahidi, MD¹²; and Shunji Takahashi, MD¹⁴

J Clin Oncol 2020;38(17):1887-96.

Effect of Trastuzumab Deruxtecan in Heavily Pretreated* HER2-Low Metastatic Breast Cancer



Clinical activity (by independent review)

ORR		
	Overall	37%
	HER2 2+	39%
	HER2 1+	36%
	ER+	40% (N = 47)
	ER-	14% (N = 7)
PFS		
	Overall	11.1 months

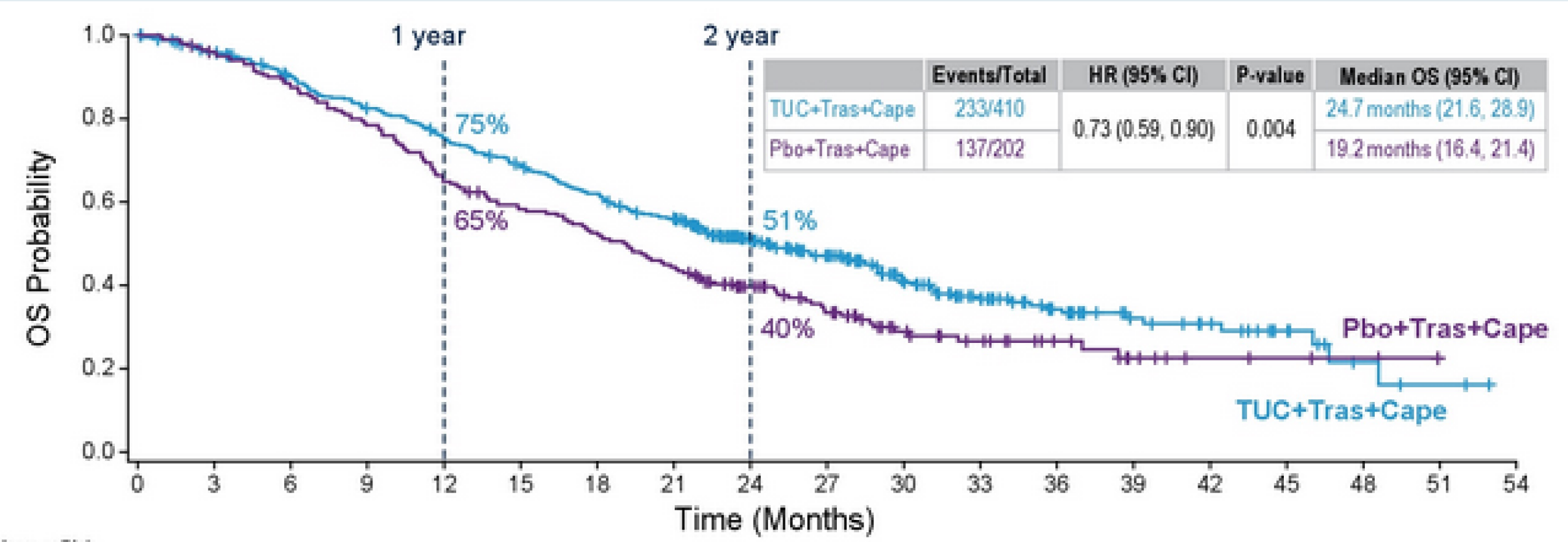
* Median of 7.5 prior regimens

Updated Results of Tucatinib versus Placebo Added to Trastuzumab and Capecitabine for Patients with Pretreated HER2+ Metastatic Breast Cancer with and without Brain Metastases (HER2CLIMB)

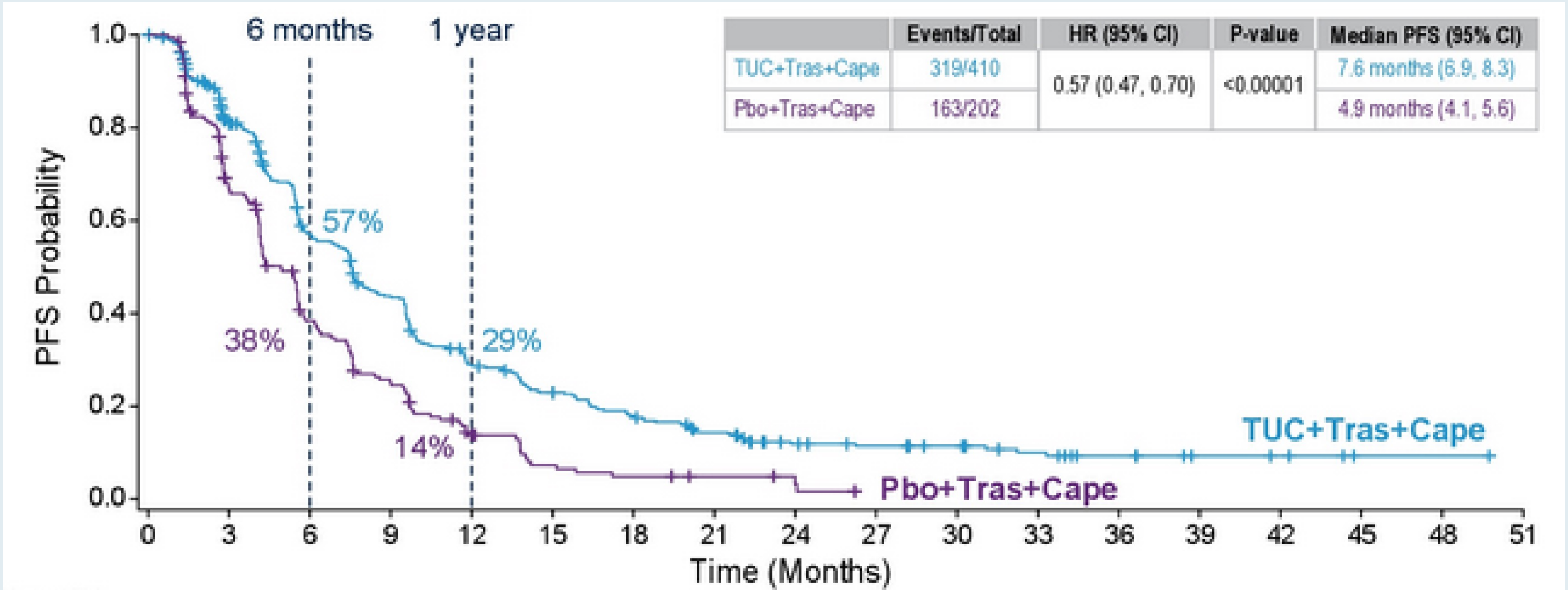
Curigliano G et al.

ASCO 2021;Abstract 1043.

HER2CLIMB: Overall Survival



HER2CLIMB: Progression-Free Survival



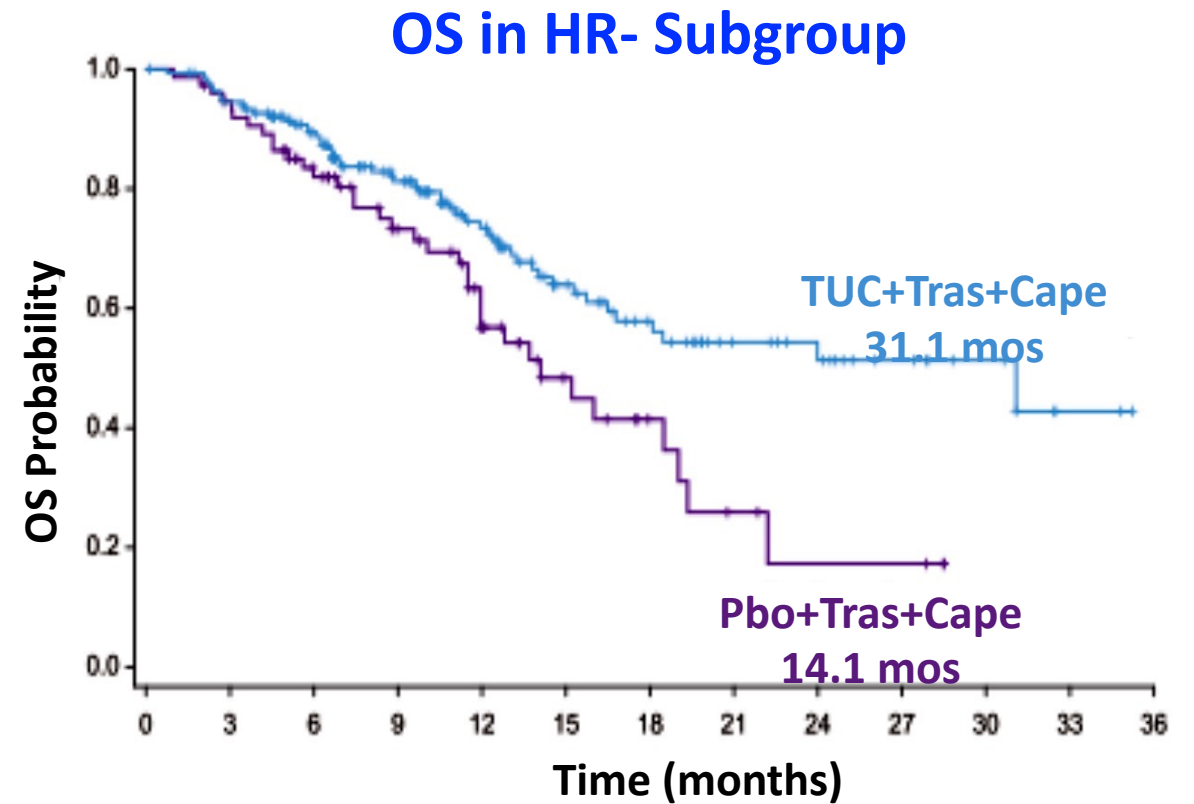
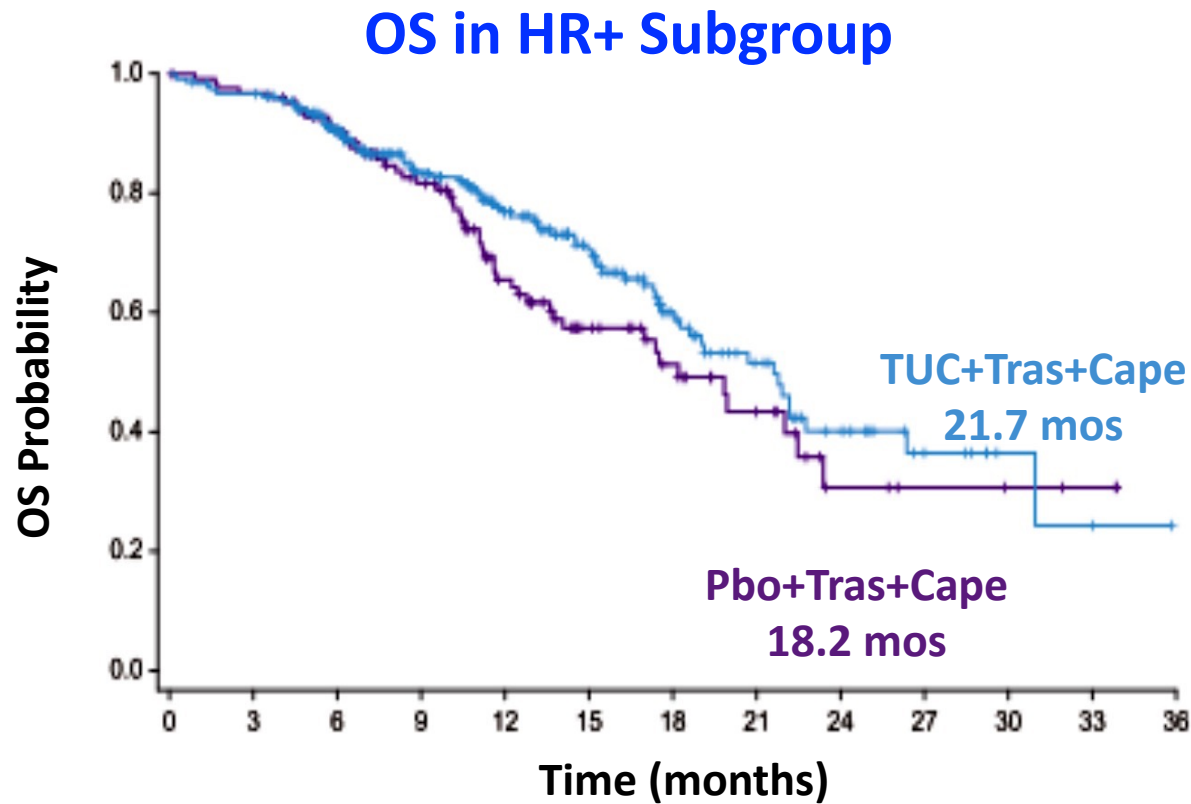
Tucatinib vs Placebo in Combination with Trastuzumab and Capecitabine for Patients with Locally Advanced Unresectable or HER2-Positive Metastatic Breast Cancer (HER2CLIMB): Outcomes by Hormone Receptor Status

Hamilton E et al.

SABCS 2020;Abstract PD3-08.

OS by HR Status in the Total Study Population

- Clinically meaningful improvement of OS was observed in patients on the tucatinib arm regardless of hormone receptor status.



HER2CLIMB: Safety Outcomes

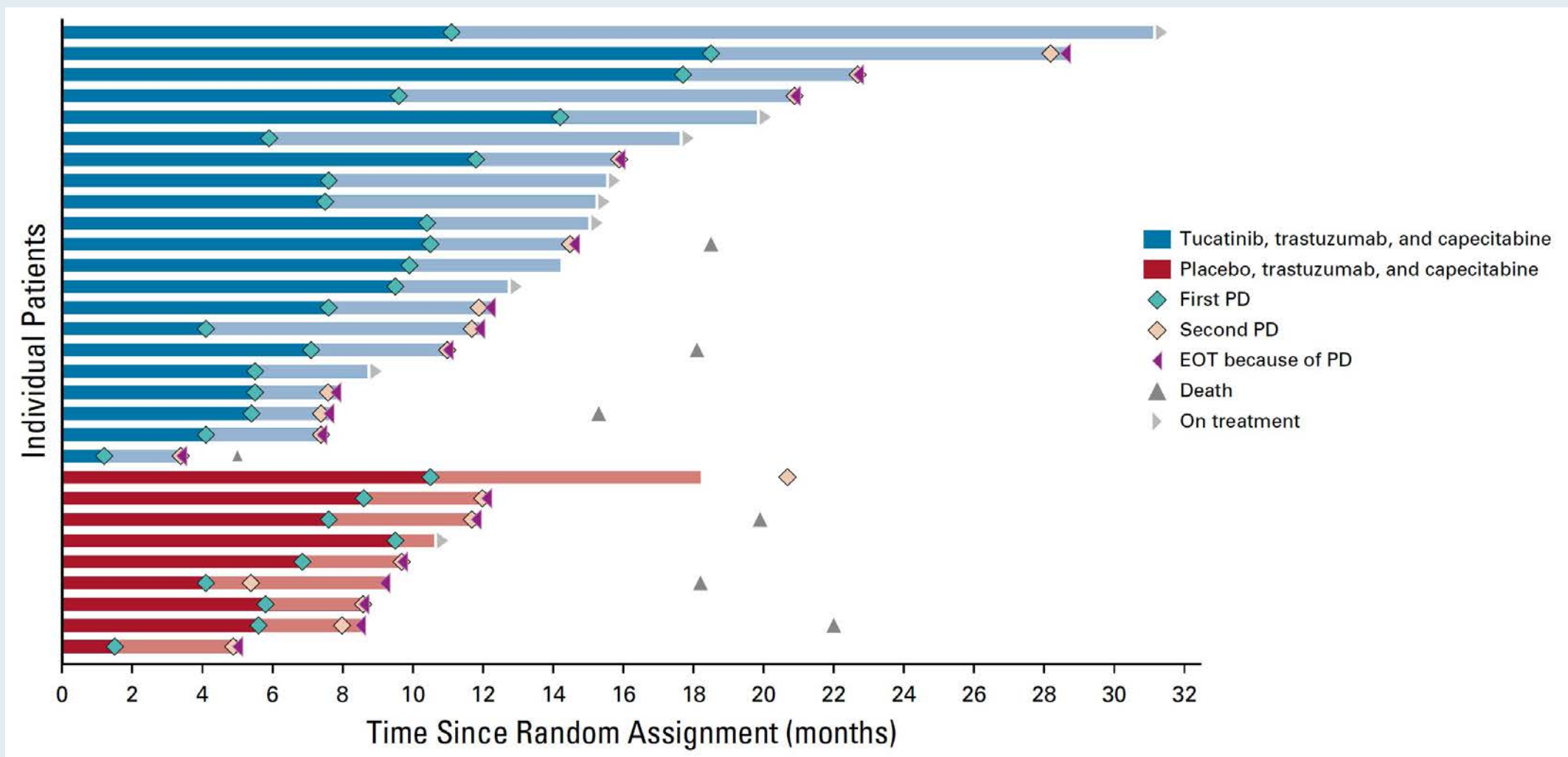
Select AE	Tucatinib (n = 404)		Placebo (n = 197)	
	Any grade	Grade ≥3	Any grade	Grade ≥3
Any	99.3%	55.2%	97.0%	48.7%
Diarrhea	80.9%	12.9%	53.3%	8.6%
PPE syndrome	63.4%	13.1%	52.8%	9.1%
Nausea	58.4%	3.7%	43.7%	3.0%
Fatigue	45.0%	4.7%	43.1%	4.1%
Vomiting	35.9%	3.0%	25.4%	3.6%
Stomatitis	25.5%	2.5%	14.2%	0.5%
Increased AST	21.3%	4.5%	11.2%	0.5%
Increased ALT	20.0%	5.4%	6.6%	0.5%

© rapid communications

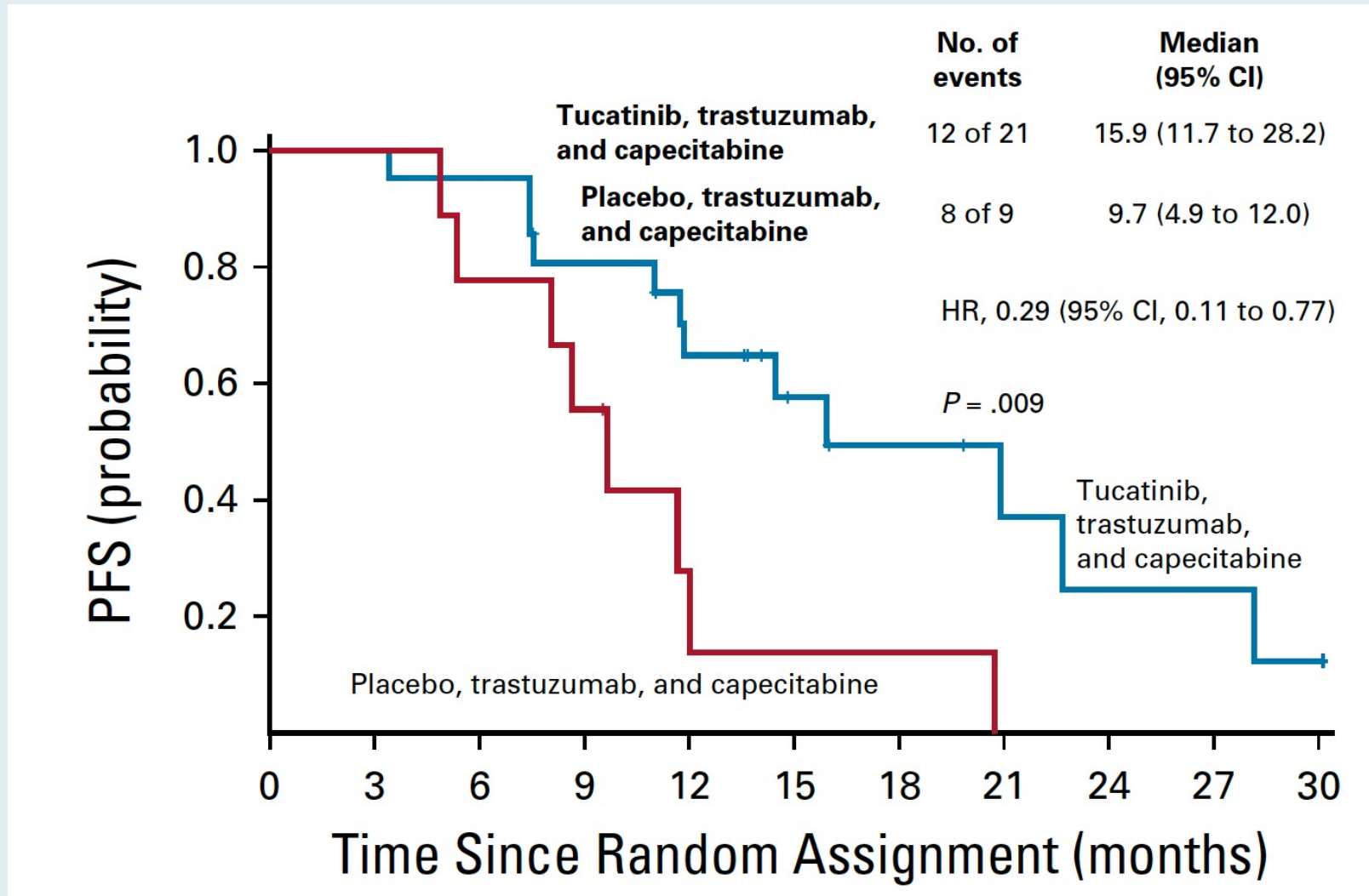
Intracranial Efficacy and Survival With Tucatinib Plus Trastuzumab and Capecitabine for Previously Treated HER2-Positive Breast Cancer With Brain Metastases in the HER2CLIMB Trial

Nancy U. Lin, MD¹; Virginia Borges, MMSc, MD²; Carey Anders, MD³; Rashmi K. Murthy, MD, MBE⁴; Elisavet Paplomata, MD⁵; Erika Hamilton, MD⁶; Sara Hurvitz, MD⁷; Sherene Loi, MD, PhD⁸; Alicia Okines, MBChB, MD⁹; Vandana Abramson, MD¹⁰; Philippe L. Bedard, MD¹¹; Mafalda Oliveira, MD, PhD¹²; Volkmar Mueller, MD¹³; Amelia Zelnak, MD¹⁴; Michael P. DiGiovanna, MD, PhD¹⁵; Thomas Bachelot, MD¹⁶; A. Jo Chien, MD¹⁷; Ruth O'Regan, MD⁵; Andrew Wardley, MBChB, MSc, MD¹⁸; Alison Conlin, MD, MPH¹⁹; David Cameron, MD, MA²⁰; Lisa Carey, MD²¹; Giuseppe Curigliano, MD, PhD²²; Karen Gelmon, MD²³; Sibylle Loibl, MD, PhD²⁴; JoAl Mayor, PharmD²⁵; Suzanne McGoldrick, MD, MPH²⁵; Xuebei An, PhD²⁵; and Eric P. Winer, MD¹

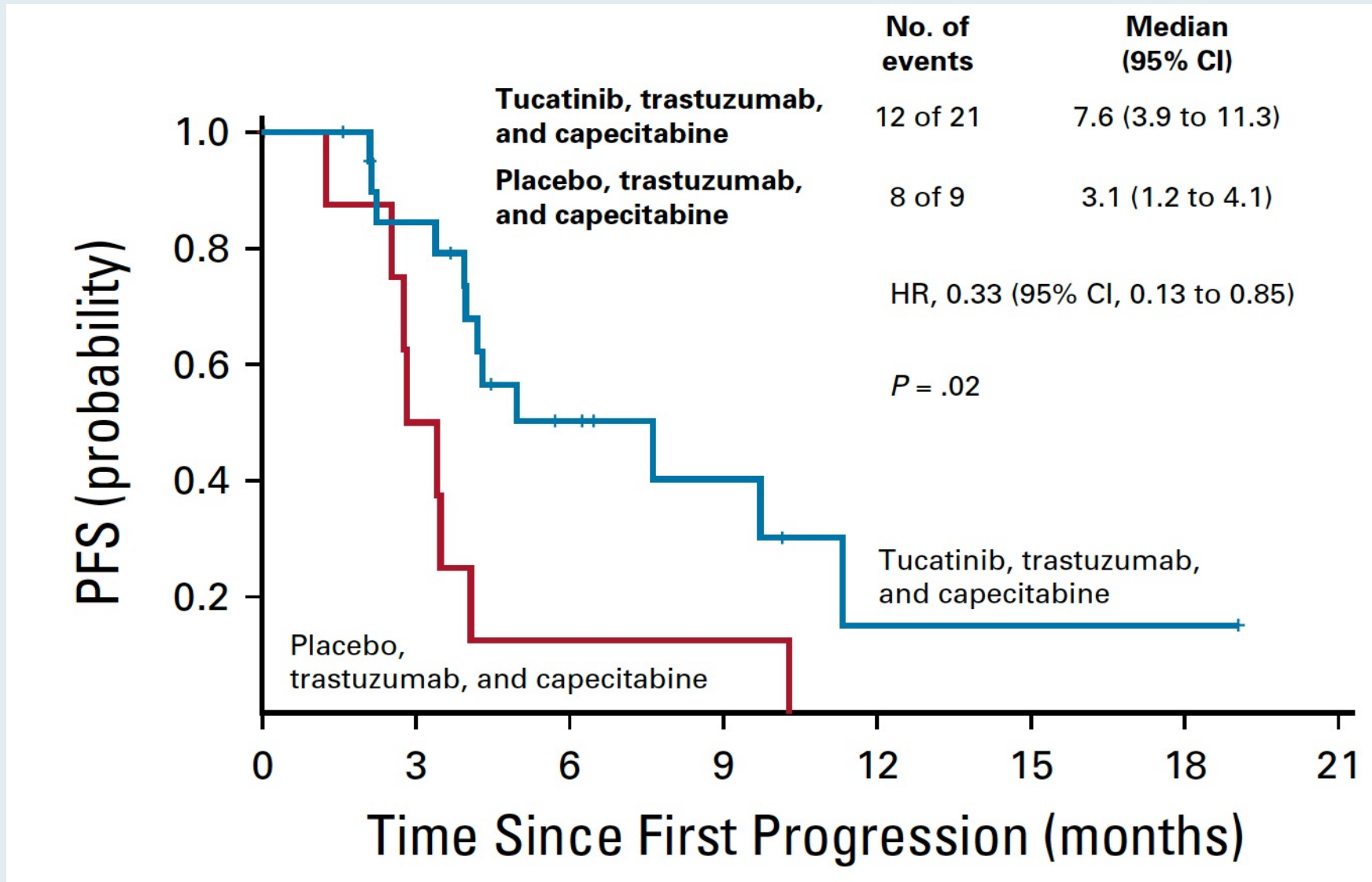
Duration of Treatment



Time from Random Assignment to Second Disease Progression by Investigator Assessment or Death



Time from First PD to Second PD by Investigator Assessment or Death



Final Overall Survival Results from the SOPHIA Study for Patients with HER2-Positive Metastatic Breast Cancer Did Not Demonstrate a Statistically Significant Advantage with Margetuximab Over Trastuzumab

Press Release – September 07, 2021

“Final overall survival (OS) results of the SOPHIA Phase 3 study in adult patients with metastatic HER2-positive breast cancer did not demonstrate a statistically significant advantage for margetuximab over trastuzumab.

The final OS analysis of the SOPHIA study was performed after 385 OS events occurred in the intent-to-treat (ITT) population. As per the study protocol, OS was defined as the number of days from randomization to the date of death (from any cause). The final OS analysis for the ITT population did not demonstrate a statistically significant advantage for margetuximab plus chemotherapy compared to that of patients who received trastuzumab plus chemotherapy (hazard ratio [HR]=0.95; 95% Confidence Interval [CI]: 0.77-1.17; P=0.62). In this overall ITT population, the median survival was 21.6 months in patients treated with margetuximab plus chemotherapy (N=266) compared to 21.9 months in patients treated with trastuzumab plus chemotherapy (N=270).

The safety profile at the time of the final OS analysis of SOPHIA was similar to what was previously reported.”



Research

JAMA Oncology | **Original Investigation**

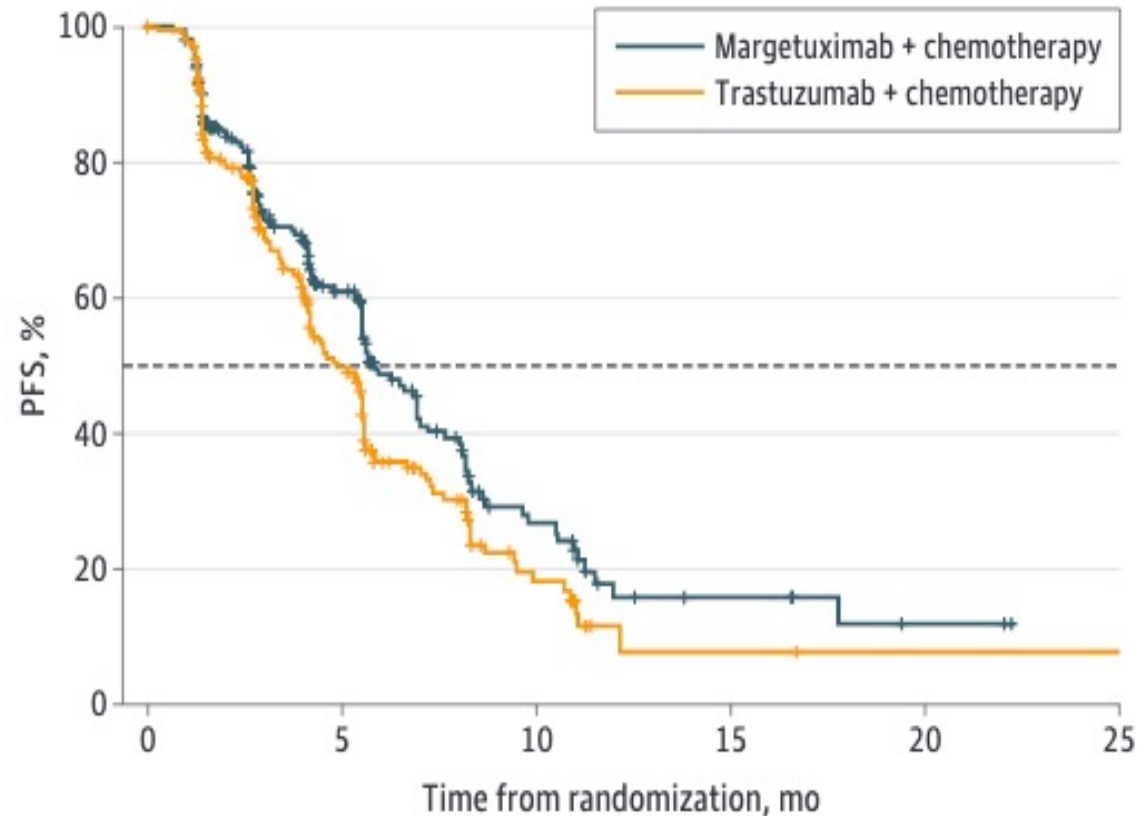
Efficacy of Margetuximab vs Trastuzumab in Patients With Pretreated ERBB2-Positive Advanced Breast Cancer

A Phase 3 Randomized Clinical Trial

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JAMA Oncol 2021;[Online ahead of print].

SOPHIA: PFS by Central Blinded Analysis (ITT Population)



	Margetuximab + chemotherapy (n = 266)	Trastuzumab + chemotherapy (n = 270)
No. of events	130	135
Median PFS (95% CI)	5.8 mo (5.52-6.97)	4.9 mo (4.17-5.59)
3-mo PFS rate	72% (65%-77%)	70% (63%-76%)
6-mo PFS rate	48% (41%-56%)	36% (28%-44%)
9-mo PFS rate	30% (22%-38%)	22% (15%-30%)

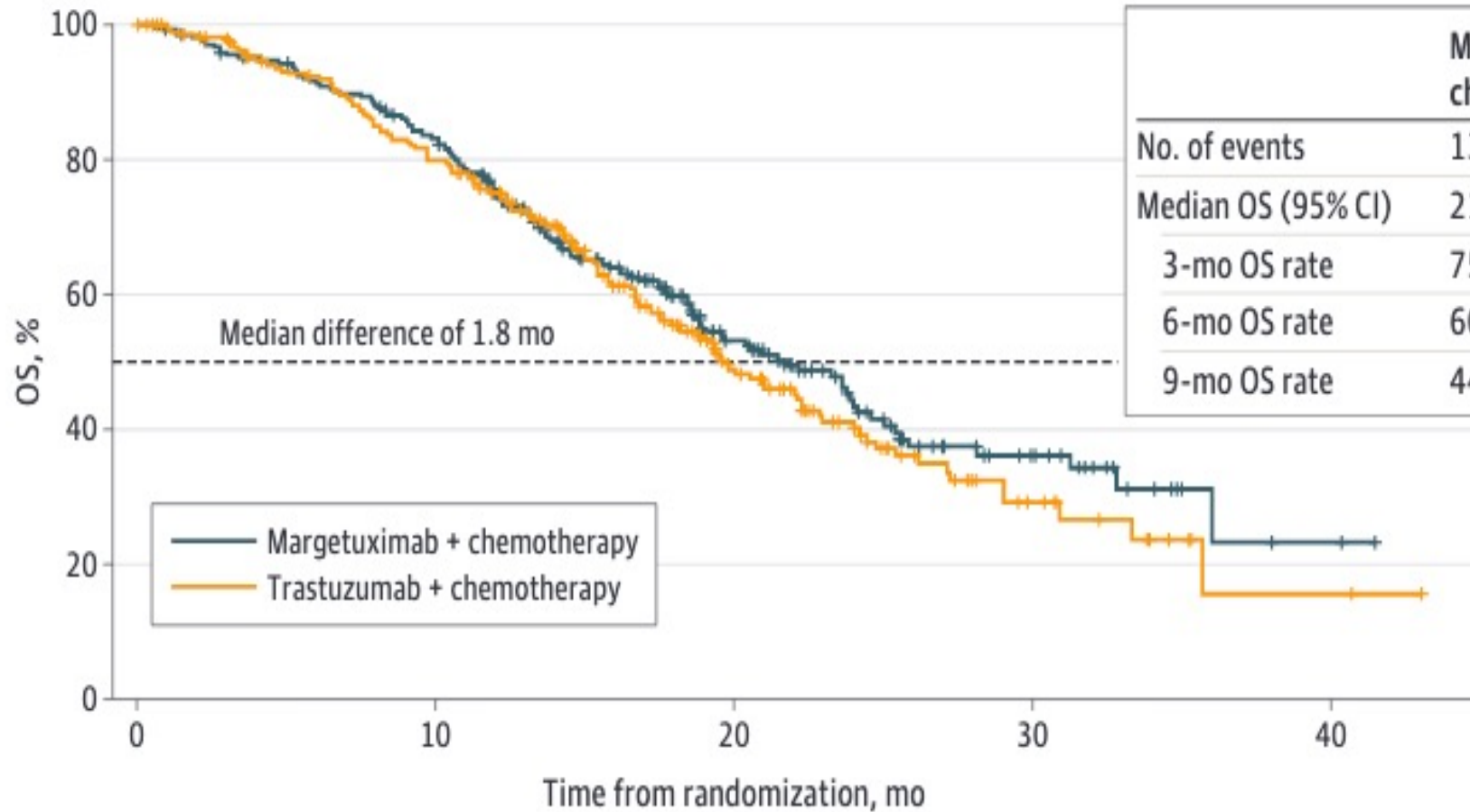
HR by stratified Cox model, 0.76 (95% CI, 0.59-0.98)

Stratified log-rank $P = .03$

24% Risk reduction of disease progression^a

Median follow-up, 2.8 mo

SOPHIA: OS Analysis (ITT Population)



	Margetuximab + chemotherapy (n = 266)	Trastuzumab + chemotherapy (n = 270)
No. of events	131	139
Median OS (95% CI)	21.6 mo (18.86-24.05)	19.8 mo (17.54-22.28)
3-mo OS rate	75% (70%-80%)	75% (70%-80%)
6-mo OS rate	60% (53%-66%)	56% (49%-62%)
9-mo OS rate	44% (36%-51%)	40% (33%-48%)

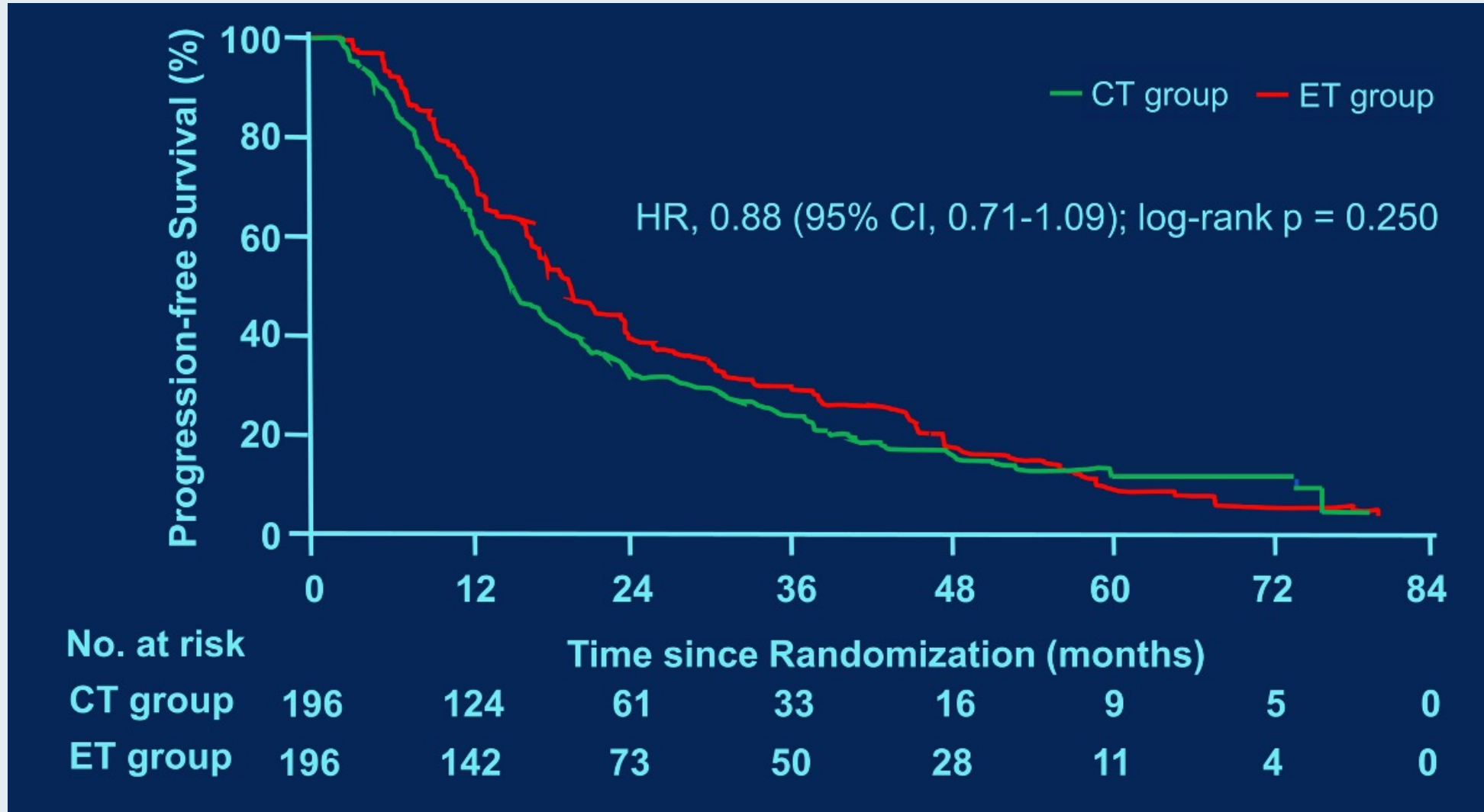
HR by stratified Cox model, 0.89 (95% CI, 0.69-1.13)
 Stratified log-rank $P = .33$
 Median follow-up, 15.6 mo

Trastuzumab plus Endocrine Therapy or Chemotherapy as First-Line Treatment for Metastatic Breast Cancer with Hormone Receptor- Positive and HER2-Positive: The SYSUCC-002 Randomized Clinical Trial

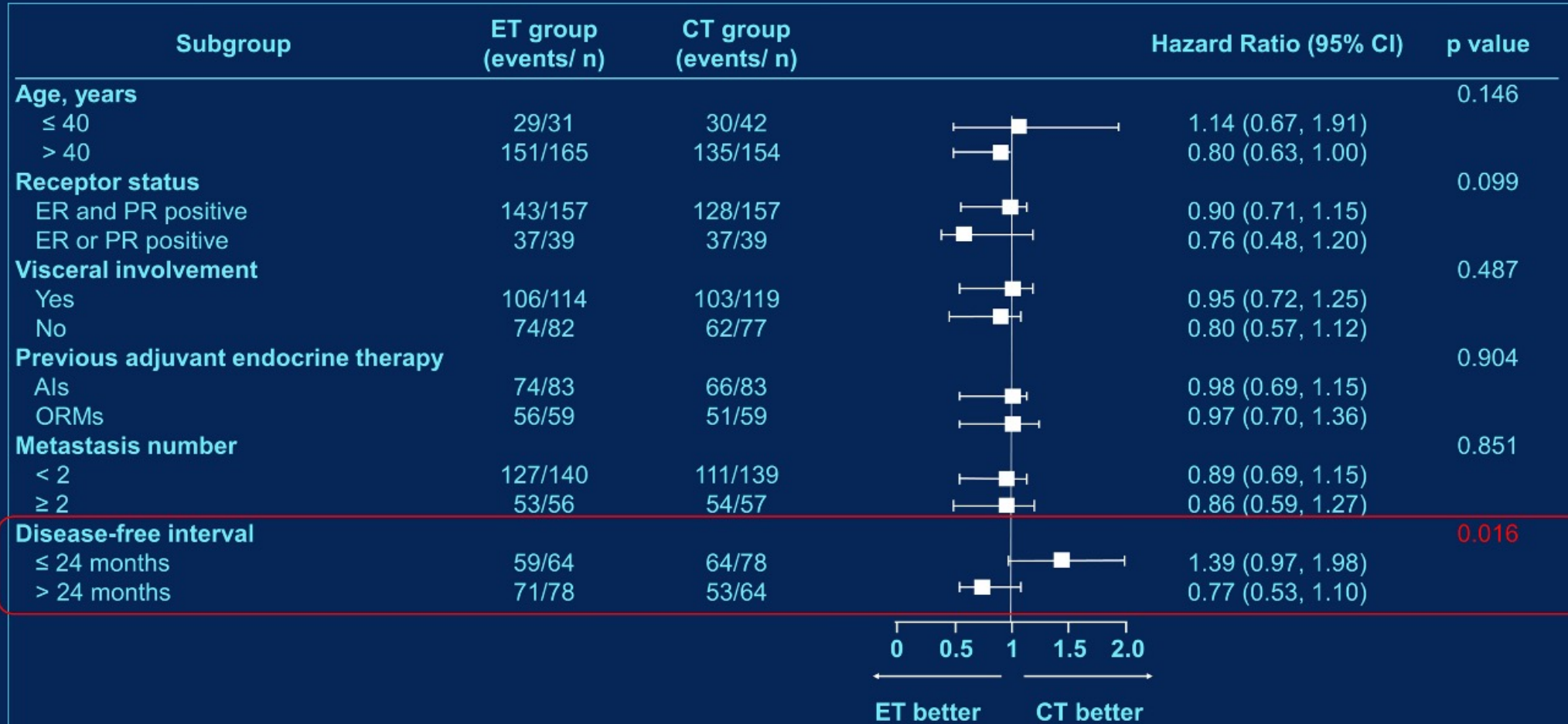
Yuan Z et al.

ASCO 2021;Abstract 1003.

SYSUCC-002: Progression-Free Survival (Primary Endpoint)



SYSUCC-002: Subgroup Analysis of PFS



Primary Outcome of the Phase III SYD985.002/TULIP Trial Comparing [vic-]Trastuzumab Duocarmazine to Physician's Choice Treatment in Patients with Pre-treated HER2-Positive Locally Advanced or Metastatic Breast Cancer

Manich E et al.

ESMO 2021;Abstract LBA15.

Conclusions: Treatment with [vic-]trastuzumab duocarmazine significantly improved PFS in comparison with standard physician's choice chemotherapy and may provide a new treatment option for patients with pre-treated locally advanced or metastatic HER2-positive breast cancer.

Select Ongoing Phase III Trials in Metastatic HER2-Positive Breast Cancer

Trial identifier	Estimated enrollment	Setting	Regimens	Estimated completion date
DESTINY-Breast09 (NCT04784715)	1,134	First line	<ul style="list-style-type: none"> Trastuzumab deruxtecan Trastuzumab deruxtecan + pertuzumab Trastuzumab + pertuzumab + taxane 	2029
HER2CLIMB-02 (NCT03975647)	460	Second line	<ul style="list-style-type: none"> T-DM1 + tucatinib Placebo + T-DM1 	2024
DESTINY-Breast02 (NCT03523585)	600	Third line	<ul style="list-style-type: none"> Trastuzumab deruxtecan Physician's choice of capecitabine/trastuzumab or capecitabine/lapatinib 	2024
DESTINY-Breast12	500	≤2 lines of therapy, presence or absence of BM	<ul style="list-style-type: none"> Trastuzumab deruxtecan 	2024

BM = brain metastases

Select Trials in Progress for HER2-Positive Breast Cancer

- ESMO 2021: 330TiP Trastuzumab deruxtecan (T-DXd; DS-8201) in HER2-positive (HER2+) and HER2-low expressing (HER-LE) metastatic breast cancer (MBC) with brain metastases (BM) and/or leptomeningeal carcinomatosis (LMC): DEBBRAH
Presenter: Marta Vaz Batista
- ESMO 2021: 329TiP KATE3 – A phase III study of trastuzumab emtansine (T-DM1) in combination with atezolizumab or placebo in patients with previously treated HER2-positive and PD-L1–positive locally advanced or metastatic breast cancer
Presenter: Sherene Loi
- ESMO 2021: 328TiP Phase III study of trastuzumab deruxtecan (T-DXd) with or without pertuzumab vs a taxane, trastuzumab and pertuzumab in first-line (1L), human epidermal growth factor receptor 2-positive (HER2+) metastatic breast cancer (mBC): DESTINY-Breast09
Presenter: Sara Tolaney

Select Trials in Progress for HER2-Positive Breast Cancer (Continued)

- ESMO 2021: 331TiP HER2CLIMB-04 – Phase II trial of tucatinib + trastuzumab deruxtecan in patients with HER2+ locally advanced or metastatic breast cancer with and without brain metastases
Presenter: Lisa Carey
- ESMO 2020: 353TiP HER2CLIMB-02 – A randomized, double-blind, phase III study of tucatinib or placebo with T-DM1 for unresectable locally advanced or metastatic HER2+ breast cancer
Presenter: Sara Hurvitz
- ASCO 2021: TPS595 Postneoadjuvant T-DM1 + tucatinib/placebo in patients with residual HER2-positive invasive breast cancer
Presenter: Ciara Catherine Maria O’Sullivan
- ASCO 2021: TPS596 eMonarchHER – A phase 3 study of abemaciclib plus standard adjuvant endocrine therapy in patients with HR+, HER2+, node-positive, high-risk early breast cancer
Presenter: Sara Tolaney

Select Trials in Progress for HER2-Positive Breast Cancer (Continued)

- ASCO 2021: TPS1099 Phase I/II study of radiation therapy followed by intrathecal trastuzumab/pertuzumab in the management of HER2+ breast leptomeningeal disease
Presenter: Kamran A Ahmed
- SABCS 2020: OT-03-01 Trastuzumab deruxtecan (T-DXd; DS-8201) vs trastuzumab emtansine (T-DM1) in high-risk patients with HER2-positive, residual invasive early breast cancer after neoadjuvant therapy: A randomized, phase 3 trial (DESTINY-Breast05)
Presenter: Charles Geyer
- SABCS 2020: OT-28-01 HER2CLIMB-02 – A randomized, double-blind, phase 3 study of tucatinib or placebo with T-DM1 for unresectable locally-advanced or metastatic HER2+ breast cancer
Presenter: Sara Hurvitz
- SABCS 2020: OT-28-03 VICKI – A Phase Ib/II, randomized, placebo-controlled, study of venetoclax plus ado-trastuzumab emtansine (T-DM1) in patients (pts) with previously treated HER2-positive locally advanced (LA) or metastatic breast cancer (MBC)
Presenter: Geoffrey Lindeman

Select Trials in Progress for HER2-Positive Breast Cancer (Continued)

- SABCS 2019: OT2-01-02 TBCRC049 – A phase II non-randomized study to assess the safety and efficacy of the combination of tucatinib and trastuzumab and capecitabine for treatment of leptomeningeal metastases in HER2 positive breast cancer
Presenter: Rashmi K Murthy

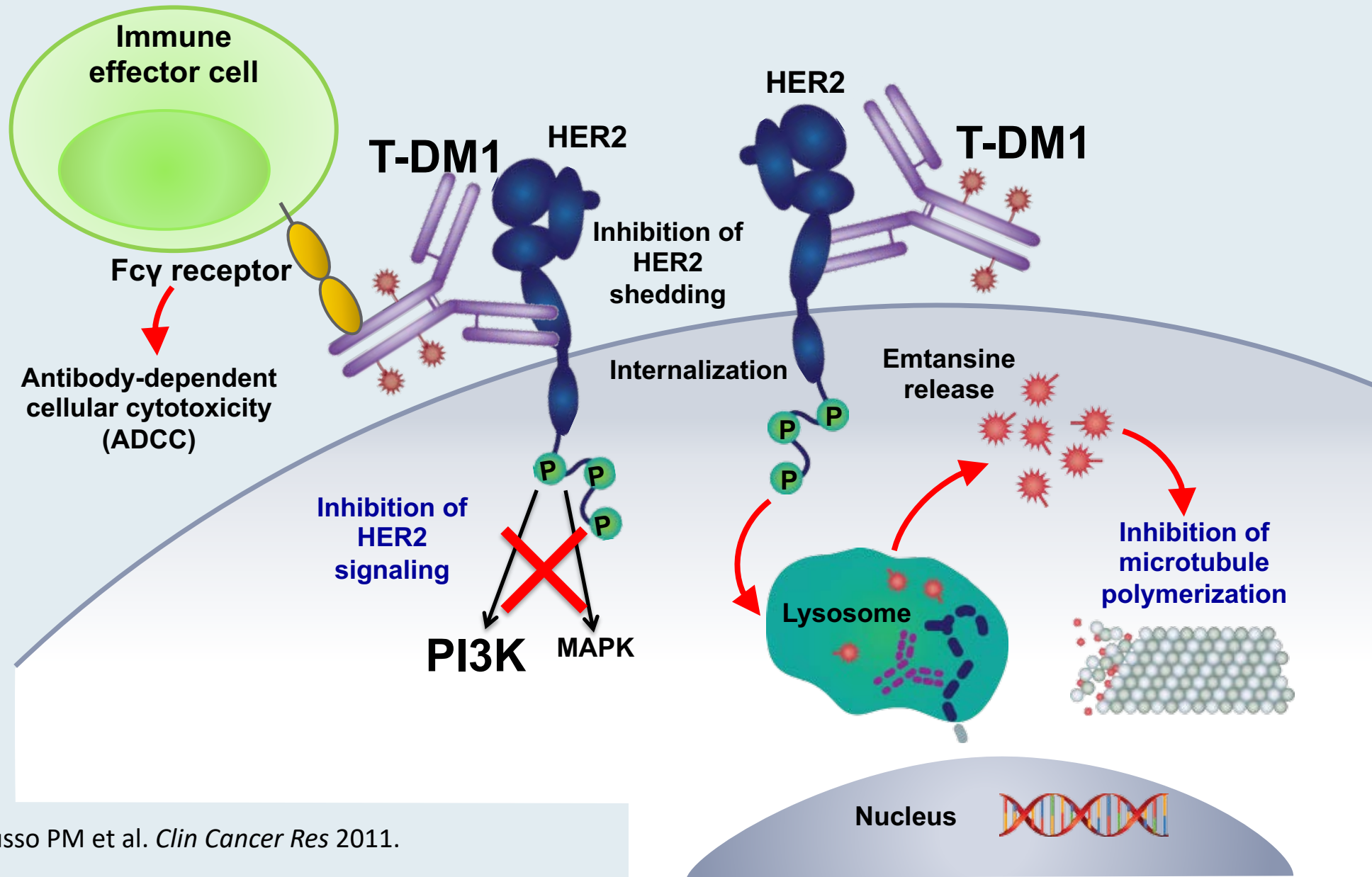
Localized HER2-Positive Breast Cancer

FDA-Approved Agents for Early-Stage HER2-Positive Breast Cancer

Agent	Setting	Pivotal trial(s)	Regimens	Year approved
Trastuzumab	Adjuvant HER2+ EBC, first line	NSABP-31 N9831 BCIRG 006 HERA	AC-T-placebo vs AC-T-H AC-T vs AC-H vs AC-T-H ACT vs ACT-H vs TC-H Observation vs trastuzumab	2006
Pertuzumab	Neoadjuvant HER2+, EBC	NeoSphere	TD vs PTD vs PT vs PD	2013
Pertuzumab	Adjuvant HER2+, EBC	APHINITY	Chemotherapy plus trastuzumab plus pertuzumab vs placebo	2017
Neratinib	Extended adjuvant treatment of HER2+ EBC	ExteNET	Placebo vs neratinib	2017
T-DM1	Adjuvant HER2+ EBC with residual disease after neoadjuvant taxane and trastuzumab-based treatment	KATHERINE	Trastuzumab vs T-DM1	2019

AC-H = doxorubicin, cyclophosphamide, and trastuzumab; AC-T, doxorubicin, cyclophosphamide, and paclitaxel; AC-T-H, doxorubicin, cyclophosphamide, paclitaxel, and trastuzumab; H, trastuzumab; PD, pertuzumab and docetaxel; PT, trastuzumab and pertuzumab; PTD, pertuzumab, trastuzumab, and docetaxel; TC, docetaxel and cyclophosphamide; TC-H, docetaxel, cyclophosphamide, and trastuzumab; TD, trastuzumab and docetaxel; THP, docetaxel, trastuzumab, and pertuzumab

Trastuzumab Emtansine (T-DM1): Mechanisms of Action



Adapted from LoRusso PM et al. *Clin Cancer Res* 2011.

ORIGINAL ARTICLE

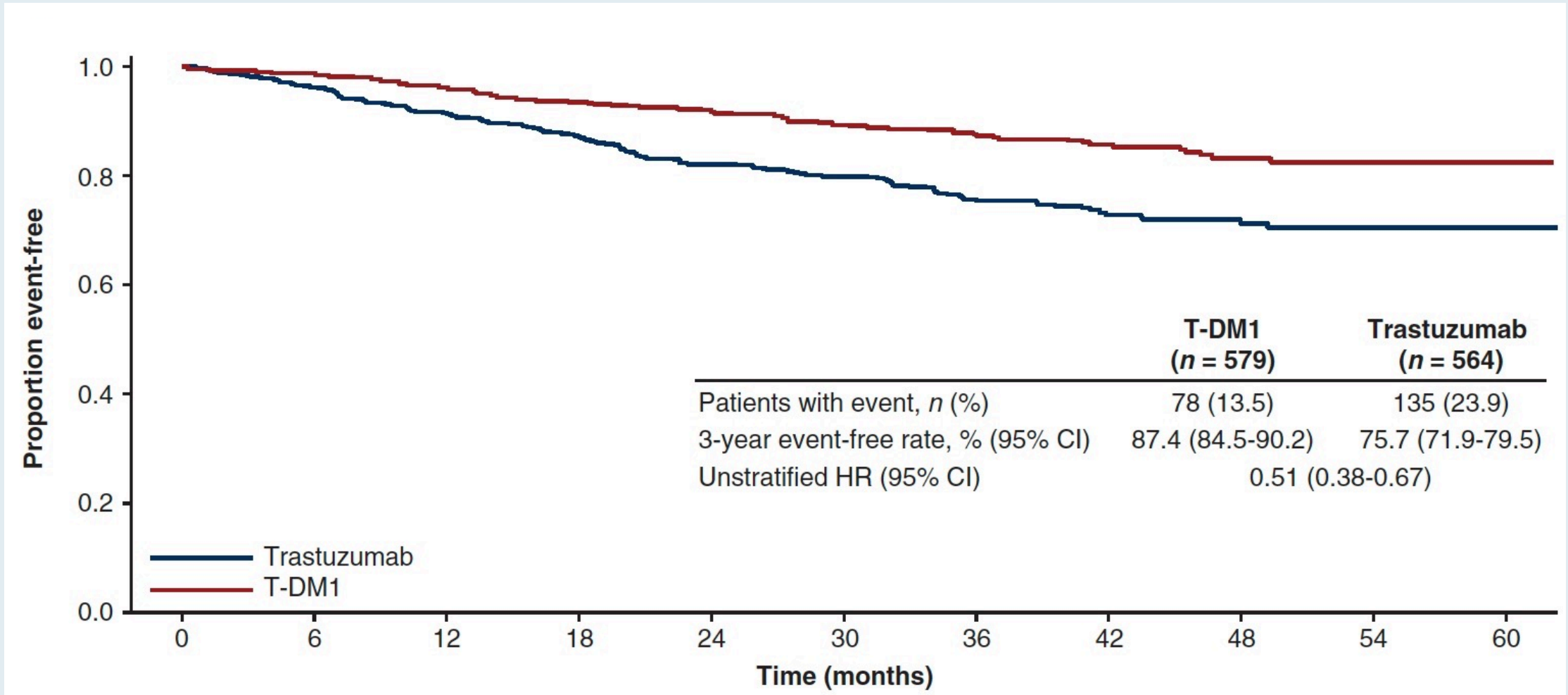
Adjuvant T-DM1 versus trastuzumab in patients with residual invasive disease after neoadjuvant therapy for HER2-positive breast cancer: subgroup analyses from KATHERINE

E. P. Mamounas^{1,2*}, M. Untch³, M. S. Mano⁴, C.-S. Huang⁵, C. E. Geyer Jr^{1,6}, G. von Minckwitz⁷, N. Wolmark^{1,8}, X. Pivot⁹, S. Kuemmel^{10,11}, M. P. DiGiovanna¹², B. Kaufman¹³, G. Kunz^{7,14}, A. K. Conlin^{1,15}, J. C. Alcedo¹⁶, T. Kuehn¹⁷, I. Wapnir^{1,18}, A. Fontana¹⁹, J. Hackmann^{7,20}, J. Polikoff^{1,21}, M. Saghatchian²², A. Brufsky^{1,23}, Y. Yang²⁴, M. Zimovjanova²⁵, T. Boulet²⁶, H. Liu²⁷, D. Tesarowski²⁸, L. H. Lam²⁸, C. Song²⁸, M. Smitt^{28,29} & S. Loibl^{7,30}

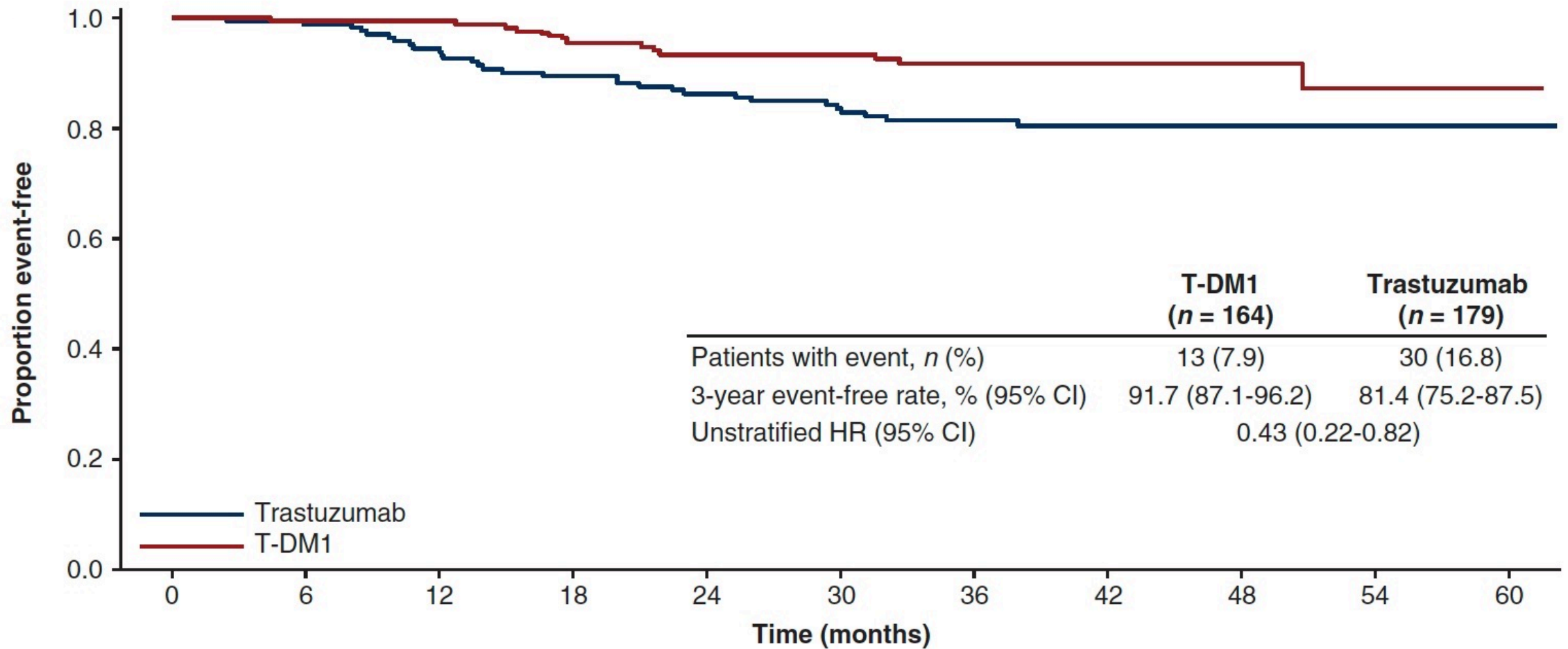
KATHERINE: Summary of Adverse Events Associated with T-DM1

Event	Trastuzumab (N = 720)	T-DM1 (N = 740)
Grade ≥3 adverse event	15.4%	25.7%
AE leading to drug discontinuation	2.1%	18.1%
Selected Grade ≥3 adverse event		
Decreased platelet count	0.3%	5.7%
Hypertension	1.2%	2.0%
Peripheral sensory neuropathy	0	1.4%
Decreased neutrophil count	0.7%	1.2%
Hypokalemia	0.1%	1.2%
Fatigue	0.1%	1.1%
Anemia	0.1%	1.1%

Time to First Invasive Disease-Free Survival Event for Patients Who Received Anthracycline-Based Neoadjuvant Therapy



Time to First Invasive Disease-Free Survival Event for Patients Who Received Non-Anthracycline-Based Neoadjuvant Therapy



Adjuvant Trastuzumab Emtansine Versus Paclitaxel in Combination With Trastuzumab for Stage I HER2-Positive Breast Cancer (ATEMPT): A Randomized Clinical Trial

Sara M. Tolaney, MD, MPH^{1,2}; Nabihah Tayob, PhD¹; Chau Dang, MD³; Denise A. Yardley, MD⁴; Steven J. Isakoff, MD, PhD⁵; Vicente Valero, MD⁶; Meredith Faggen, MD¹; Therese Mulvey, MD⁵; Ron Bose, MD, PhD⁷; Jiani Hu, MSc¹; Douglas Weckstein, MD¹; Antonio C. Wolff, MD⁸; Katherine Reeder-Hayes, MD, MBA, MSc⁹; Hope S. Rugo, MD¹⁰; Bhuvanewari Ramaswamy, MD¹¹; Dan Zuckerman, MD¹²; Lowell Hart, MD¹³; Vijayakrishna K. Gadi, MD, PhD¹⁴; Michael Constantine, MD¹; Kit Cheng, MD¹⁵; Frederick Briccetti, MD¹; Bryan Schneider, MD¹⁶; Audrey Merrill Garrett, MD¹⁷; Kelly Marcom, MD¹⁸; Kathy Albain, MD¹⁹; Patricia DeFusco, MD²⁰; Nadine Tung, MD^{2,21}; Blair Ardman, MD²²; Rita Nanda, MD²³; Rachel C. Jankowitz, MD²⁴; Mothaffar Rimawi, MD²⁵; Vandana Abramson, MD²⁶; Paula R. Pohlmann, MD, PhD, MSc²⁷; Catherine Van Poznak, MD²⁸; Andres Forero-Torres, MD²⁹; Minetta Liu, MD³⁰; Kathryn Ruddy, MD³⁰; Yue Zheng, MSc¹; Shoshana M. Rosenberg, ScD, MPH^{1,2}; Richard D. Gelber, PhD^{1,2}; Lorenzo Trippa, PhD^{1,2}; William Barry, PhD¹; Michelle DeMeo, BS¹; Harold Burstein, MD, PhD^{1,2}; Ann Partridge, MD, MPH^{1,2}; Eric P. Winer, MD^{1,2}; and Ian Krop, MD, PhD^{1,2}

J Clin Oncol 2021;[Online ahead of print]

ATEMPT: Invasive Disease-Free Survival (iDFS) and Recurrence-Free Interval (RFI)

Outcome	T-DM1 (n = 383)	TH (n = 114)
Three-year iDFS	97.8%	93.4%
Three-year RFI	99.2%	94.3%

ATEMPT: Clinically Relevant Toxicity

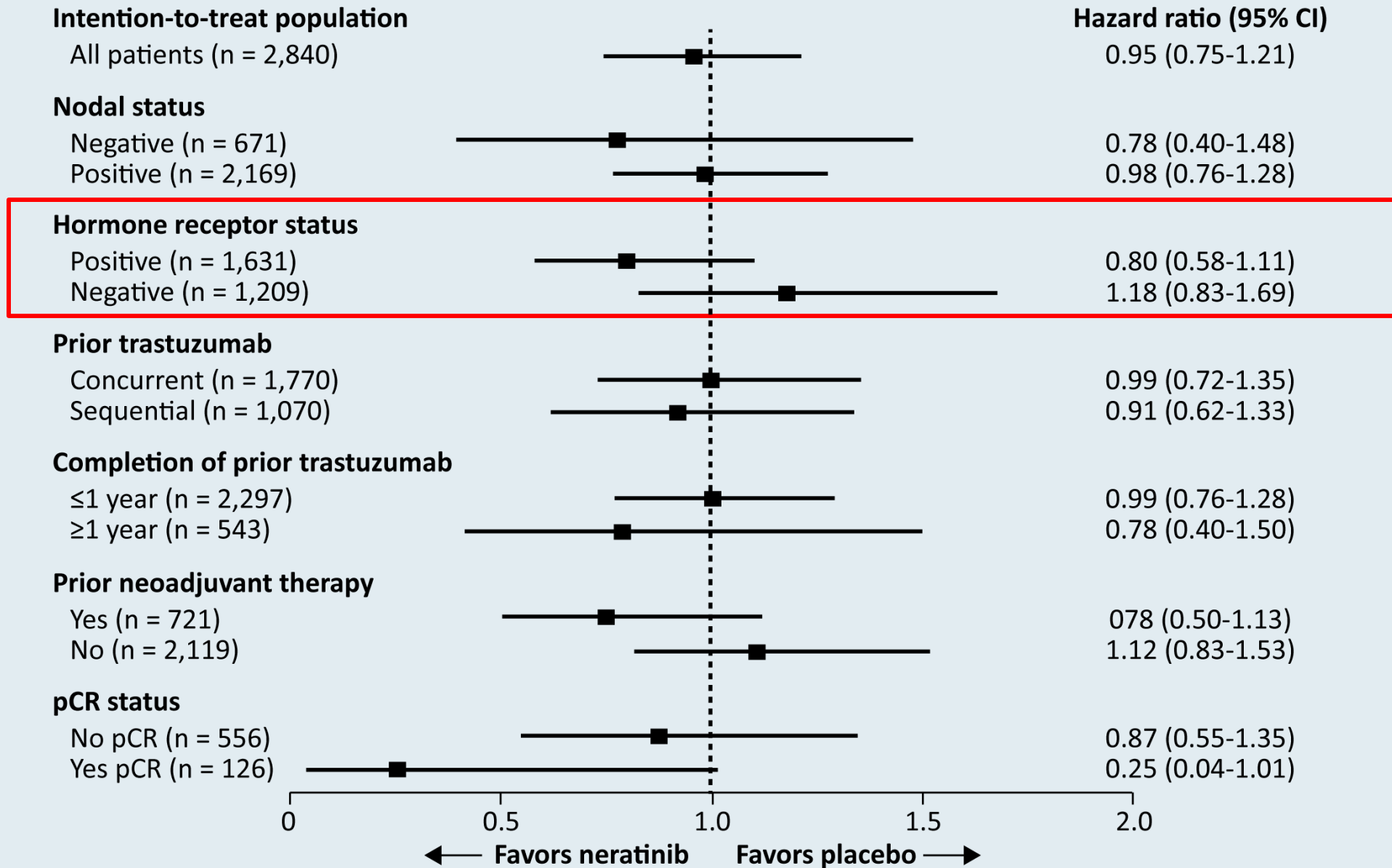
Clinically Relevant Toxicity	T-DM1 (n = 383)	TH (n = 114)
Grade ≥ 3 nonhematologic toxicity	9%	11%
Grade ≥ 2 neurotoxicity	11%	23%
Grade ≥ 4 hematologic toxicity	1%	0%
Febrile neutropenia	0%	2%
Any toxicity requiring dose delay	28%	26%
Any toxicity requiring early discontinuation	17%	6%
Total	46%	47%

Continued Efficacy of Neratinib in Patients with HER2-Positive Early-Stage Breast Cancer: Final Overall Survival Analysis from the Randomized Phase 3 ExteNET Trial

Holmes FA et al.

SABCS 2020;Abstract PD3-03.

ExteNET: Final Overall Survival Analysis



ExteNET: Cumulative Incidence of CNS Recurrences

Population or subgroup	Events, n		Cumulative incidence of CNS recurrences, % (95% CI)	
	Neratinib	Placebo	Neratinib	Placebo
Intention-to-treat population (n = 2,840)	16	23	1.3 (0.8-2.1)	1.8 (1.2-2.7)
HR+/\leq1-year population (EU indication) (n = 1,334)	4	12	0.7 (0.2-1.7)	2.1 (1.1-3.5)
Prior neoadjuvant therapy (n = 1,334)				
No (n = 980)	3	6	0.7 (0.2-2.0)	1.5 (0.6-3.0)
Yes (n = 354)	1	6	0.7 (0.1-3.3)	3.7 (1.5-7.4)
pCR status (n = 354)				
No (n = 295)	1	5	0.8 (0.1-4.0)	3.6 (1.3-7.8)
Yes (n = 38)	0	1	0 (NE)	5.0 (0.3-21.2)

ExteNET: CNS Disease-Free Survival at 5 Years

Population or subgroup	Events, n		Kaplan-Meier estimate at 5 years %, (95% CI)		Hazard ratio
	Neratinib	Placebo	Neratinib	Placebo	
Intention-to-treat population (n = 2,840)	29	42	97.5 (96.4-98.3)	96.4 (95.2-97.4)	0.73
HR+/\leq1-year population (EU indication) (n = 1,334)	9	23	98.4 (96.8-99.1)	95.7 (93.6-97.2)	0.41
Prior neoadjuvant therapy (n = 1,334)					
No (n = 980)	7	10	98.2 (96.3-99.2)	97.5 (95.3-98.6)	0.70
Yes (n = 354)	2	13	98.7 (94.8-99.7)	91.2 (85.1-94.8)	0.18
pCR status (n = 354)					
No (n = 295)	2	10	98.4 (93.6-99.6)	92.0 (85.6-95.7)	0.24
Yes (n = 38)	0	3	100 (100-100)	81.9 (53.1-93.9)	0

CONTROL Trial: Strategies to Improve Neratinib Tolerability

Background: Neratinib is approved for extended adjuvant therapy in HER2-positive BC

- Neratinib poorly tolerated in ExteNET
 - Discontinuation rate: 17%
 - Grade 3 diarrhea: 40%

Objective: Improve GI tolerability of neratinib

Methods: Sequential single arm interventions in patients treated with adjuvant therapy

- Cohort 1 (L): Loperamide (n = 137)
- Cohort 2 (BL): Budesonide + loperamide (n = 64)
- Cohort 3 (CL or CL-PRN): Colestipol + loperamide (n = 136) or colestipol + as needed loperamide (n = 104)
- Cohort 4 (DE): Neratinib dose escalation; ongoing (n = 60)

Treatment-Emergent Diarrhea in the ExteNET and CONTROL Studies

Outcome	ExteNET (n = 1408)	L (n = 137)	BL (n = 64)	CL (n = 136)	CL-PRN (n = 104)	DE (n = 60)
Treatment-emergent diarrhea incidence, n (%)						
No diarrhea	65 (5)	28 (20)	9 (14)	23 (17)	5 (5)	1 (2)
Grade 1	323 (23)	33 (24)	16 (25)	38 (28)	34 (33)	25 (42)
Grade 2	458 (33)	34 (25)	21 (33)	47 (35)	32 (31)	25 (42)
Grade 3	561 (40)	42 (31)	18 (28)	28 (21)	33 (32)	9 (15)
Grade 4	1 (<1)	0	0	0	0	0
Action taken, n (%)						
Dose hold	477 (34)	20 (15)	12 (19)	22 (16)	15 (14)	7 (12)
Dose reduction	372 (26)	10 (7)	3 (5)	10 (7)	12 (12)	2 (3)
Discontinuation	237 (17)	28 (20)	5 (8)	5 (4)	8 (8)	2 (3)
Hospitalization	20 (1)	2 (1)	0	0	0	0

Select Ongoing Trials in Early-Stage HER2-Positive Breast Cancer

Trial identifier	Phase	Setting	Regimens	Estimated completion date
CompassHER2 pCR (NCT04266249)	II	Neoadjuvant and adjuvant	<ul style="list-style-type: none"> • Preoperative chemotherapy + trastuzumab/pertuzumab • <i>If pCR</i> → postoperative trastuzumab/pertuzumab • <i>If residual disease</i> → postoperative T-DM1 or T-DM1 + tucatinib 	2023
DESTINY-Breast05 (NCT04622319)	III	High-risk, residual disease after neoadjuvant chemotherapy	<ul style="list-style-type: none"> • Trastuzumab deruxtecan • T-DM1 	2027

Key Considerations in the Optimal Clinical Care of Patients with Small Cell Lung Cancer

A CME/MOC-Accredited Virtual Event

Thursday, November 4, 2021

5:00 PM – 6:00 PM ET

Faculty

Anne Chiang, MD, PhD

David R Spigel, MD

Moderator

Neil Love, MD

Thank you for joining us!

***CME and MOC credit information will be emailed
to each participant within 5 business days.***